

ORIGINAL PATENT APPLICATION BASED ON:

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**ELECTROLUMINESCENT DEVICES INCLUDING CONJUGATED
POLYMERS CONTAINING AN AZOLE STRUCTURE**

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**ELECTROLUMINESCENT DEVICES INCLUDING CONJUGATED
POLYMERS CONTAINING AN AZOLE STRUCTURE**

FIELD OF THE INVENTION

5 The present invention relates to conjugated polymers containing an
azole structure for use in electroluminescent (EL) devices.

BACKGROUND OF THE INVENTION

Electroluminescent devices are opto-electronic devices where light
emission is produced in response to an electrical current through the device. The
physical model for EL is the radiative recombination of electrons and holes. The
10 term light emitting diode (LED) is commonly used to describe an EL device
where the current-voltage behavior is non-linear, meaning that the current through
the EL device is dependent on the polarity of the voltage applied to the EL device.
Both organic and inorganic materials have been used for the fabrication of LEDs.
Inorganic materials such as ZnS/Sn, Ga/Bs, Ga/As have been used in
15 semiconductor lasers, small area displays, LED lamps, etc. However, the
drawbacks of inorganic materials include difficulties to process and to obtain large
surface areas and efficient blue light.

Organic materials, which includes both small molecules and
polymeric materials, offer several advantages over inorganic materials for LEDs,
20 such as simpler manufacturing, low operating voltages, the possibility of
producing large area and full-color displays. Conjugated polymers such as
poly(phenylvinylene) (PPV) were first introduced as EL materials by Burroughes
et al in 1990 (Burroughes, J. H. et al, *Nature* 1990, 347, 539-41). Tremendous
progress has been made since then to improve the stability, efficiency, and
25 durability of polymeric LEDs (Bernius, M. T. et al, *Adv. Mater.* 2000, 12, 1737).
Organic LED (OLED) represents an alternative to the well-established display
technologies based on cathode-ray tubes and liquid crystal displays (LCDs),
especially for large area displays. OLED has been demonstrated to be brighter,
thinner, lighter, and faster than LCDs. Moreover it requires less power to operate,
30 offers higher contrast and wide viewing angle (>165 degree), and has great

potential to be cheaper to manufacture, especially the polymer-based LEDs (PLED).

The OLED technology has stimulated intensive research activities across all disciplines. Currently, great efforts in materials research have been
5 focused on efficient novel materials for full-color flexible displays. Full-color displays require three basic colors, red, green and blue, and flexible substrates require low temperature and easy processing of the organic materials. PLED devices show great promise in meeting both requirements, since the emission color can be tailored by modulation of the chemical structures and the solution
10 processing allows for micro-patterning of the fine multicolor pixels via inkjet printing technique. However, processable, stable, and efficient blue light emitting organic materials are still highly desirable to meet the challenge. Blue light requires a wide energy band. With blue light emitting polymers as primary materials, it is possible to produce other colors by a downhill energy transfer
15 process. For instance, a green or red EL emission can be obtained by doping a blue EL host material with a small amount of green or red luminescent material.

Furthermore, efficient EL devices require balanced charge injection and transport. The turn-on voltage depends on the energy barriers for charge injection. The mobility of the injected charges in organic polymers is low
20 compared to that in inorganic semiconductors, which might increase the recombination time for the electrons and holes in the polymer. Moreover, most EL polymers transport preferentially either electrons or holes that leads to the recombination zone close to the electrodes and quenching of excitons. This is one of the main sources of reduced efficiency. Many research efforts have been
25 focused on the design of polymers with improved charge injection and transporting.

It is known that imine nitrogens (C=N) are electron-deficient and are capable of transporting electrons. Oxadiazoles, pyridines, quinolines, quinoxazolines, triazines, and bithiozoles all process electron transporting
30 properties. Electron transporting small organic molecules such as 2-(4-biphenyl)-5-(4-tert-butylphenyl)-1,3,4-oxadiazole (PBD) have been dispersed in

inert polymers or blended with EL polymers to improve electron transporting capability (Zhang, C. et al. *Synth. Met.* 1995, 72, 185; Berggren, M. et al. *Adv. Mater.* 1995, 7, 900). However, the incompatibility in general leads to phase separation or recrystallization especially at higher temperature during device operation. Oxadiazoles containing polymers have been intensively investigated. A number of polymers containing oxadiazole moiety have been reported by several research groups (Yang, Y. et al. *Chem. Mater.* 1995, 7, 1568). However, most of these polymers do not meet the requirement of low driving voltage or improved efficiency.

SUMMARY OF THE INVENTION

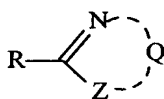
It is an object of the present invention to provide novel highly efficient luminescent materials useful in emissive layers.

It is another object of the present invention to provide wide energy band gap luminescent materials.

It is another object of the present invention to provide novel charge transport materials useful in electroluminescent devices.

These objects are achieved in an electroluminescent device, comprising:

- a) a spaced-apart anode and cathode; and
- b) an organic layer disposed between the spaced-apart anode and cathode and including a polymer having an azole structure represented by formula (I)



wherein:

Z is O, NR', or S;

Q represents atoms necessary to complete a hetero ring with N and Z;

R is a substituent and selected from hydrogen, or alkyl, or alkenyl, or alkynyl, or alkoxy wherein the alkyl, alkenyl, alkynyl or alkoxy can have from 1 to 40 carbon atoms; or aryl from 6 to 60 carbon atoms; or heteroaryl from 4 to 60

carbons; or F, or Cl, or Br; or a cyano group; or a nitro group; or atoms coupled to N or Z to complete a fused aromatic or heteroaromatic ring; and

R' is hydrogen, or alkyl, or alkenyl, or alkynyl of from 1 to 40 carbon atoms wherein the alkyl, alkenyl, alkynyl or alkoxy can have from 1 to 40 carbon atoms; aryl from 6 to 60 carbon atoms; or heteroaryl from 4 to 60 carbons; or F, or Cl, or Br.

The present invention provides organic luminescent materials with a number of advantages that include excellent solubility and thermal stability, high efficiency, and good charge transport capability.

The present invention includes a polymer having a particular azole structure that itself is emissive and can be used in an emissive layer of an electroluminescent device, and it also can be used as a host, particularly for phosphorescent dopants. This azole structure also is useful in an electron transport layer.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates in cross-section a basic structure of an EL device;
FIG. 2 illustrates the absorption (AB) and photoluminescence (PL) spectra of polymer 43 in solution and thin film;

FIG. 3 illustrates the EL spectra of the two EL devices fabricated from polymer 43: ITO/PEDOT/polymer 43/CsF/Mg:Ag and ITO/PEDOT/polymer 43 + 4 wt% Ir(PPY)₃/CsF/Mg:Ag; and

FIG. 4 illustrates the voltage-current density of the two EL devices fabricated from polymer 43 and polymer 43 doped with Ir(PPY)₃.

DETAILED DESCRIPTION OF THE INVENTION

It has been discovered that small molecule azole compounds are useful as emitting layer or electron transport layer in efficient EL devices as disclosed in commonly-assigned U.S. Patent Nos. 5,645,948 and 5,766,779. The present invention provides highly efficient organic light-emitting materials that include a conjugated polymer having an azole structure with good electron transport capability, excellent solubility, thermal stability, and improved efficiency. A conjugated polymer is one that possesses a delocalized pi-electron

system along the backbone. The azole structure is represented by formula (I), wherein:

Z is O, NR', or S;

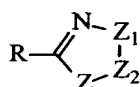
Q represents atoms necessary to complete a hetero ring with N and Z and, if possible, R and Z, or R and Q may be bound to each other to form a ring;

R is a substituent and selected from hydrogen, or alkyl, or alkenyl, or alkynyl, or alkoxy of from 1 to 40 carbon atoms; aryl or substituted aryl of from 6 to 60 carbon atoms; or heteroaryl or substituted heteroaryl of from 4 to 60 carbons; or F, or Cl, or Br; or a cyano group; or a nitro group; or atoms necessary to complete a fused aromatic or heteroaromatic ring; and

R' is hydrogen, or alkyl, or alkenyl, or alkynyl of from 1 to 40 carbon atoms; aryl or substituted aryl of from 6 to 60 carbon atoms; or heteroaryl or substituted heteroaryl of from 4 to 60 carbons; or F, or Cl, or Br; or atoms necessary to complete a fused aromatic or heteroaromatic ring.

For example, R and R' are independently hydrogen, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, t-butyl, pentyl, hexyl, ethylhexyl, heptyl, octyl, nonyl, decyl, dodecyl, hexyadecyl, cyclohexyl, cyclopentyl, methoxy, ethoxy, butoxy, hexyloxy, ethylhexyloxy, methoxyethoxyethyl, methoxyethylhexyloxyethyl, phenyl, tolyl, naphthyl, xylene, anthracene, phenanthrene, phenylmethylenepheryl, benzyl, phenoxy, pyridyl, thiophenyl, furyl, or quinolinyl. R and R' are groups that form fused aromatic or heteroaromatic rings such as naphthalene, anthracene, perylene, phenanthrene, pyrene, tetracene, pentacene, triphenylene, and benzo[a]pyrene. Preferably, R and R' are hydrogen, t-butyl, hexyl, 2-ethylhexyl, octyl, 3,7-dimethyloctyl, decyl, heptyl, phenyl, 2-ethylhexyloxy, or 4-methoxyphenyl; diphenylamino, (4-diphenylamino)phenyl; R and R' are groups that forms fused aromatic rings such as naphthalene, anthracene, perylene, pyrene, phenanthrene, or tetracene.

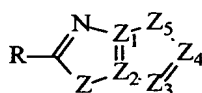
In some embodiments Q represents atoms necessary to complete a five-member hetero ring with N and Z, the azole of structure formula (I) is represented by formula (Ia), wherein:



(Ia)

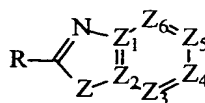
5 Z_1 and Z_2 are the same or different, and are selected from CR', O, NR', or S, and R' is defined above.

In preferred embodiments, when Z_1 and Z_2 are atoms necessary to complete a fused five- or six-member aromatic or heteroaromatic ring, the azole of structure formula (I) is represented by formula (Ib1) and (Ib2), wherein:



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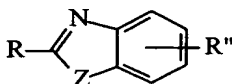
(Ib1)



(Ib2)

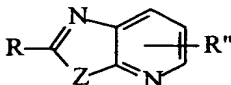
15 $Z_1, Z_2, Z_3, Z_4, Z_5,$ and Z_6 are the same or different, and $Z_3, Z_4, Z_5,$ and Z_6 are defined as the same as $Z_1,$ and Z_2 noted above.

In more preferred embodiments, when Z_1 and Z_2 are CR' or NR' to complete a fused six-member aromatic or heteroaromatic ring, the azole of structure formula (I) is represented by formula (Ic), (Id), and (Ie), wherein:



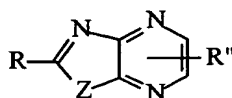
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(Ic)



(Id)

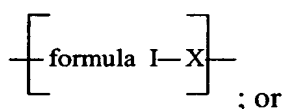
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(Ie)

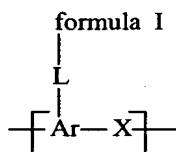
R'' is defined as the same as R noted above.

Conjugated polymers including the azole structure are represented
 5 by repeating units of formula (II) that includes the azole structure as part of the
 polymer main chain and repeating units of formula (III) which includes the azole
 structure as part of the polymer side chain.



(II)

10



(III)

wherein:

X is a conjugated group of 2 to 60 carbon atoms;

15 Ar is an aryl or substituted aryl group containing 6 to 60 carbon atoms; or
 heteroaryl or substituted heteroaryl containing 4 to 60 carbon atoms, and at least
 one or more N, S, or O atoms; and

L is a direct bond between formula (I) and Ar or a carbon linking group
 having 1 to 40 carbon atoms or a non-carbon linking group having 0 to 40 carbon
 20 atoms that includes an alkyl, alkenyl, alkynyl, aryl, or heteroaryl; L may also
 contains O, N, S, F, Cl, or Br, or Si atoms;

Incorporating X into the polymers comprising the azole structure
 represented by formula (II), and (III) can further improves solubility, or electron
 or hole transporting mobility, or finely tune the emission color.

25 The present invention includes a conjugated polymer having a particular
 azole structure that itself is emissive and can be used in an emissive layer of an

electroluminescent device. This azole structure also is useful in an electron transport layer.

5 X is a conjugated group of 2-60 carbon atoms that includes vinylenes, ethynylenes, arylenes, heteroarylenes, arylene vinylenes, heteroarylene vinylenes, and combinations thereof;

For example, arylenes contain 6 to 60 carbon atoms which include phenyl, biphenyl, naphthyl, anthracene, fluorene, phenanthrene, spirophe-
nyl, spirofluorene, perylene, or pyrene groups;

10 Heteroarylenes contain 4 to 60 carbon atoms which include pyridine, thiophene, pyrrole, bithiophene, furan, benzofuran, benzimidazole, benzoxazole, quinoxaline, phenylquinoline, diphenyloxadizaole, or carbazole;

Vinylenes, ethynylenes, arylenes, heteroarylenes, arylene vinylenes, and heteroarylene vinylenes may be substituted or unsubstituted. The substituents include but are not limited to alkyl or alkoxy groups containing 1 to 40 carbon
15 atoms, aryl groups containing 6 to 60 carbon atoms; or heteroaryl groups containing 4 to 60 carbons; or F, Cl, or Br; or a cyano group; or a nitro group.

X can be divided into the following groups.

Group I:

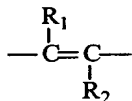
X is a vinylene, or ethynylene group of formula (IV):



wherein:

W contains 2 to 40 carbon atoms, may also contains O, N, S, F, Cl, or Br, or Si atoms.

25 The following structures constitute specific examples of formula (IV)



wherein: R₁ and R₂ are individually hydrogen, or alkyl of 1 to 40 carbon atoms, or aryl containing 6 to 60 carbon atoms; or heteroaryl containing 4 to 60 carbons; or F, Cl, or Br; or a cyano group; or a nitro group; or

30



Group II:

X is a group containing two aryl or heteroaryl groups connected by a linking group L_1 of formula (V):

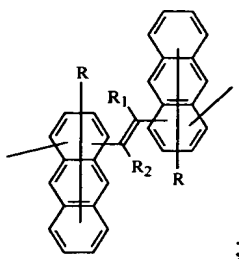
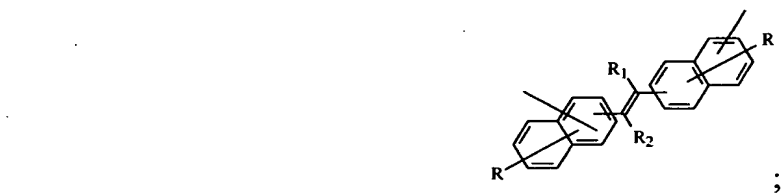
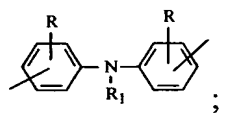


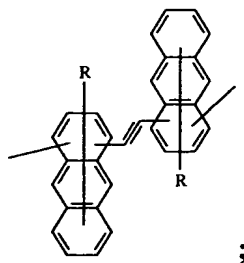
wherein:

Ar_1 and Ar_2 are aryl groups containing 6 to 60 carbon atoms, or heteroaryl groups containing 4 to 60 carbon atoms, and at least one or more N, S, or O atoms; and

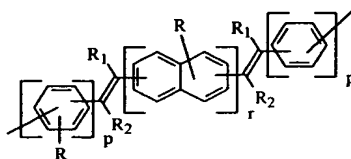
L_1 is a linking groups containing 0 to 40 carbon atoms, may contain N, Si, O, Cl, F, Br, or S atoms.

The following structures constitute specific examples of formula (V)



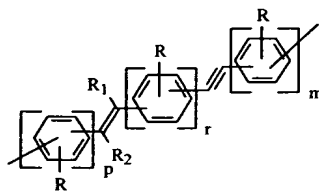


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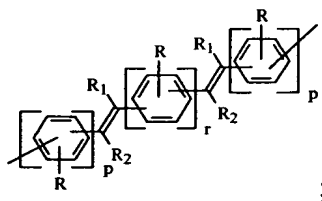


5

wherein: p and r are integers from 1 to 4;

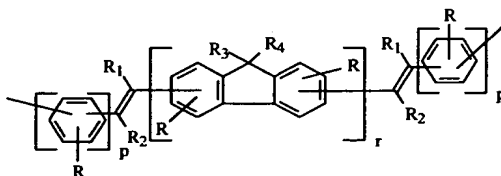


wherein: m is an integer from 1 to 4;



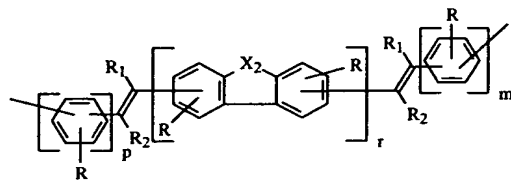
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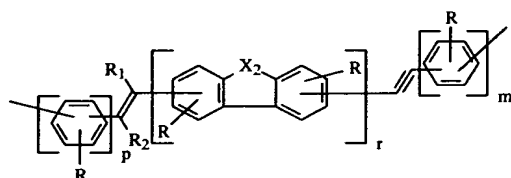


wherein: R₃ and R₄ are substituents each being individually hydrogen, or alkyl, or alkenyl, or alkynyl, or alkoxy of from 1 to 40 carbon atoms; aryl or substituted

aryl of from 6 to 60 carbon atoms; or heteroaryl or substituted heteroaryl of from 4 to 60 carbons; or F, Cl, or Br; or a cyano group; or a nitro group;

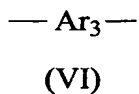


5 wherein: X_2 is S, Se, or O atom, SiR_2 , or N-R; or



Group III:

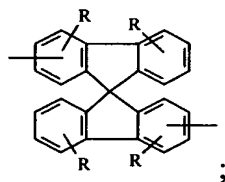
10 X is an aryl or heteroaryl group of formula (VI):



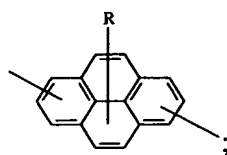
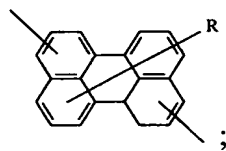
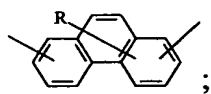
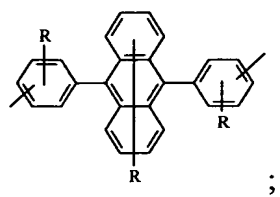
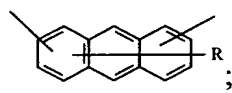
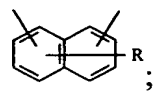
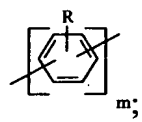
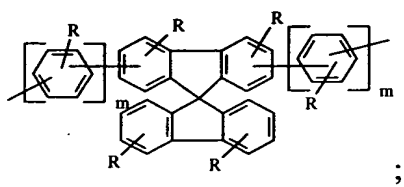
wherein:

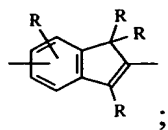
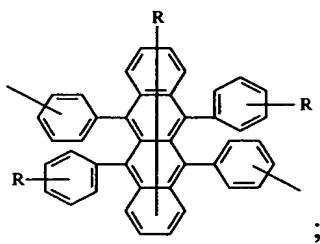
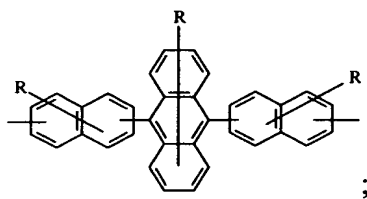
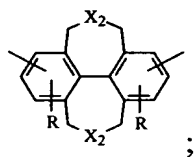
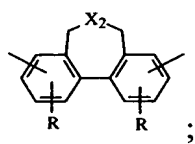
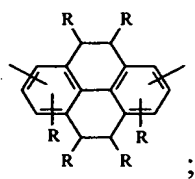
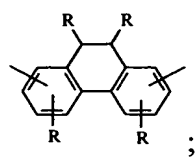
Ar₃ is an aryl or substituted aryl group containing 6 to 60 carbon atoms; or
 15 heteroaryl or substituted heteroaryl containing 4 to 60 carbon atoms, and at least one or more N, S, or O atoms;

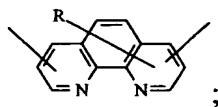
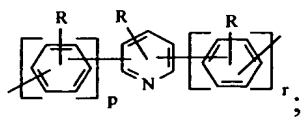
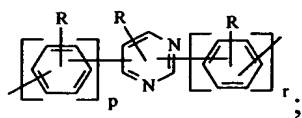
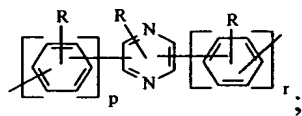
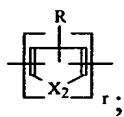
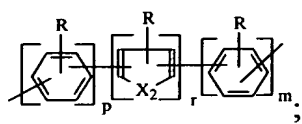
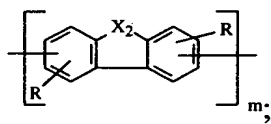
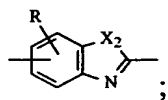
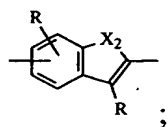
The following structures constitute specific examples of formula (VI)

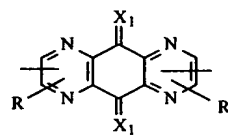
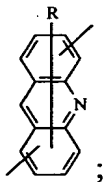
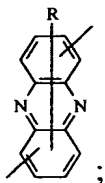
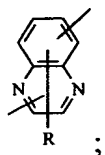
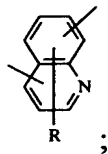
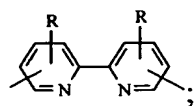


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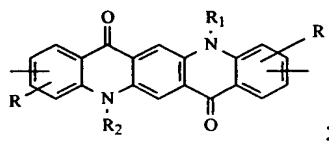


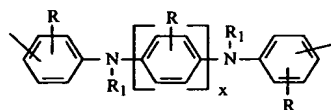
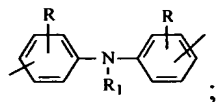
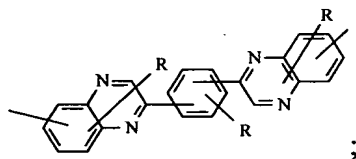
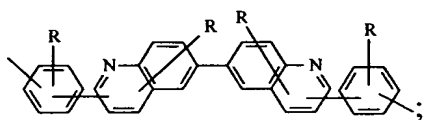
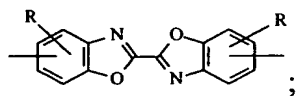
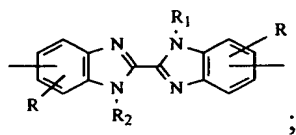
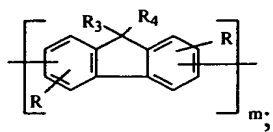




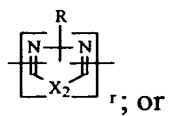


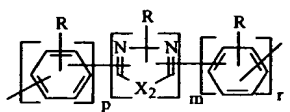
wherein: X_1 is an O atom or two cyano groups;





x is an integer of 1 to 3;



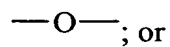


X can be one or the combination of more than one of the above groups.

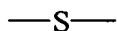
- 5 L is a direct bond between formula (I) and Ar or a carbon linking group having 1 to 40 carbon atoms or a non-carbon linking group having 0 to 40 carbon atoms. When L is a linking group, it includes but are not limited to the following groups:

10 Group I:

L is an ether, or thioether linking group:

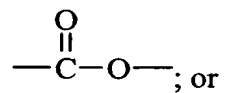


15

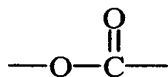


Group II:

L is an ester linking group:



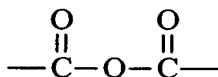
20



Group III:

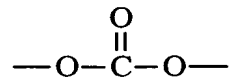
L is an anhydride linking group:

25



Group IV:

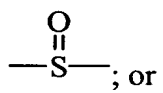
L is a carbonate linking group:



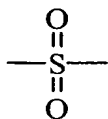
5

Group V:

L is a sulfone or sulfine linking group:

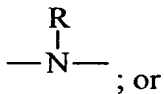


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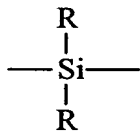


Group VI:

L is an amino or silo linking group:



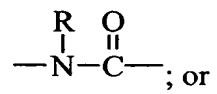
15

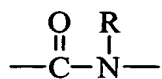


Group VII:

L is an amide linking group:

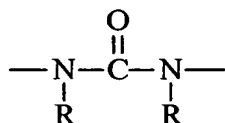
20





Group VIII:

L is a urea linking group:



5

Group IX:

L is an alkyl, alkenyl, or alkynyl group:

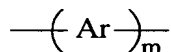


10 wherein:

Y contains 1 to 40 carbon atoms, may also contains O, N, S, F, Cl, Br, or Si atoms.

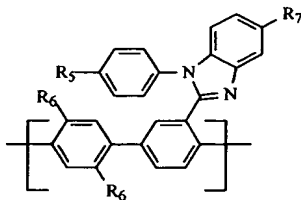
Group X:

15 L is an aryl or heteroaryl linking group:



L can be one or the combination of more than one of the above groups.

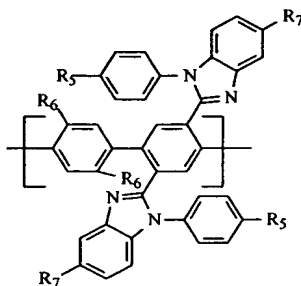
20 The following molecular structures constitute specific examples of preferred polymers satisfying the requirement of this invention:



polymer 1 $\text{R}_5 = \text{R}_6 = \text{n-hexyl}$, $\text{R}_7 = 2\text{-ethylhexyl}$

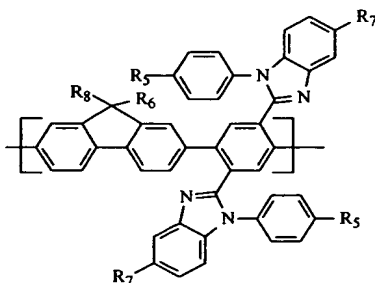
polymer 2 $\text{R}_5 = \text{H}$, $\text{R}_6 = \text{R}_7 = 3,7\text{-dimethyloctyl}$

- polymer 3 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = H$, $R_7 = t\text{-butyl}$
polymer 4 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = n\text{-decyl}$, $R_7 = H$
polymer 5 $R_5 = n\text{-hexyl}$, $R_6 = R_7 = H$
polymer 6 $R_5 = R_6 = n\text{-hexyloxy}$, $R_7 = H$
5 polymer 7 $R_5 = R_6 = R_7 = n\text{-hexyloxy}$



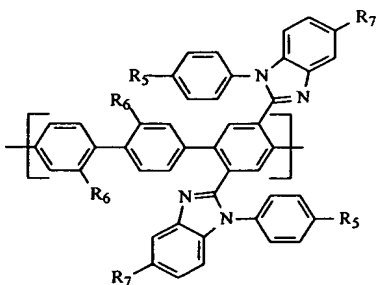
- polymer 8 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = 2\text{-ethylhexyl}$
polymer 9 $R_5 = H$, $R_6 = R_7 = 3,7\text{-dimethyloctyl}$
10 polymer 10 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = H$, $R_7 = t\text{-butyl}$
polymer 11 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = n\text{-decyl}$, $R_7 = H$
polymer 12 $R_5 = n\text{-hexyl}$, $R_6 = R_7 = H$
polymer 13 $R_5 = R_6 = n\text{-hexyloxy}$, $R_7 = H$
polymer 14 $R_5 = R_6 = R_7 = n\text{-hexyloxy}$

15



- polymer 15 $R_5 = R_6 = R_7 = R_8 = n\text{-hexyl}$
polymer 16 $R_5 = R_7 = H$, $R_6 = R_8 = 3,7\text{-dimethyloctyl}$
polymer 17 $R_5 = R_7 = H$, $R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_8 = n\text{-hexyl}$
20 polymer 18 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = R_8 = n\text{-decyl}$, $R_7 = H$

polymer 19 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = \text{n-hexyl}$, $R_8 = \text{n-octyl}$



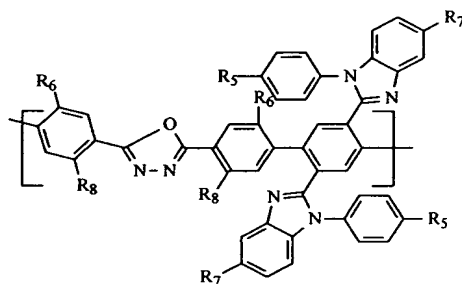
polymer 20 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

5 polymer 21 $R_5 = \text{H}$, $R_6 = R_7 = 3,7\text{-dimethyloctyl}$

polymer 22 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$, $R_7 = \text{t-butyl}$

polymer 23 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = \text{n-decyl}$, $R_7 = \text{H}$

polymer 24 $R_5 = \text{n-hexyl}$, $R_6 = R_7 = \text{H}$



10

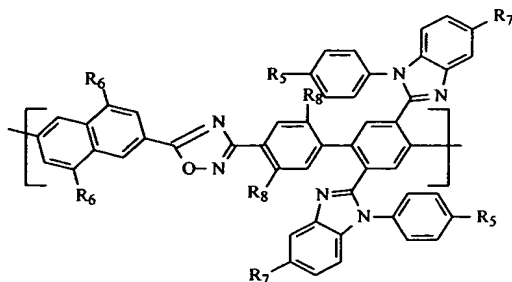
polymer 25 $R_5 = R_6 = R_7 = R_8 = \text{n-hexyl}$

polymer 26 $R_5 = \text{n-hexyl}$, $R_7 = \text{H}$, $R_6 = R_8 = 3,7\text{-dimethyloctyloxy}$

polymer 27 $R_5 = R_7 = \text{H}$, $R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_8 = \text{H}$

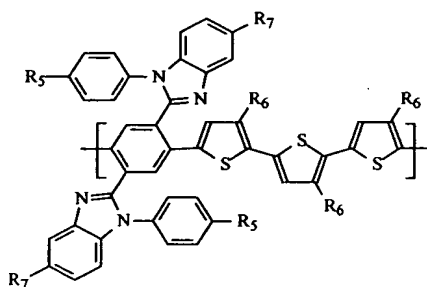
polymer 28 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = \text{n-decyl}$, $R_7 = R_8 = \text{H}$

15

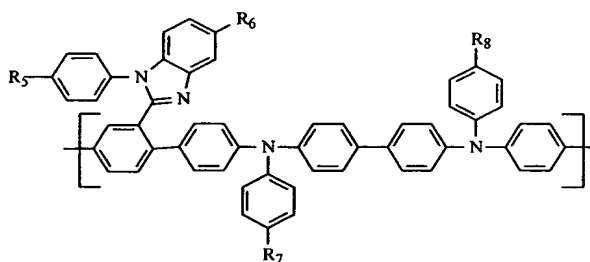


polymer 29 $R_5 = R_6 = R_7 = R_8 = \text{n-hexyl}$

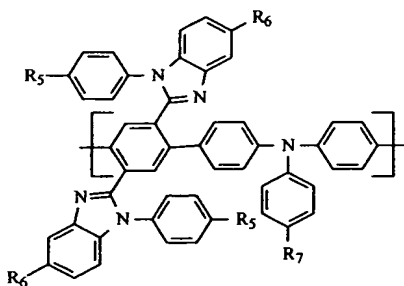
- polymer 30 $R_5 = \text{n-hexyl}, R_7 = \text{H}, R_6 = R_8 = 3,7\text{-dimethyloctyloxy}$
 polymer 31 $R_5 = R_7 = \text{H}, R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_8 = \text{H}$
 polymer 32 $R_5 = \text{n-hexyloxy}, R_6 = \text{n-decyl}, R_7 = R_8 = \text{H}$



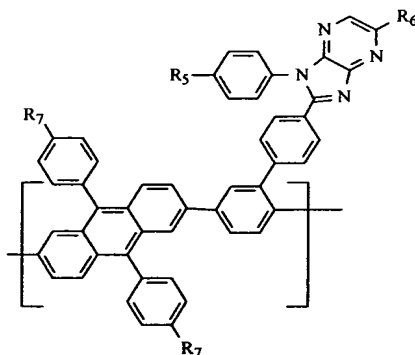
- 5
 polymer 33 $R_5 = R_6 = \text{n-hexyl}, R_7 = 2\text{-ethylhexyl}$
 polymer 34 $R_5 = \text{H}, R_6 = R_7 = 3,7\text{-dimethyloctyl}$
 polymer 35 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = 2\text{-ethylhexyl}, R_7 = \text{H}$
 10
 polymer 36 $R_5 = R_6 = \text{n-hexyl}, R_7 = \text{H}$
 polymer 37 $R_5 = R_7 = \text{n-hexyloxy}, R_6 = 2\text{-ethylhexyl}$



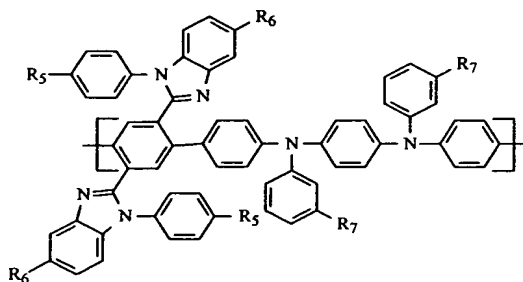
- polymer 38 $R_5 = R_6 = \text{n-hexyl}, R_7 = R_8 = 2\text{-ethylhexyl}$
 15
 polymer 39 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = R_8 = \text{t-butyl}$
 polymer 40 $R_5 = R_7 = R_8 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = \text{H}$
 polymer 41 $R_5 = R_6 = \text{n-hexyl}, R_7 = R_8 = (4\text{-carbazole})\text{phenyl}$
 polymer 42 $R_5 = \text{n-hexyloxy}, R_6 = \text{H}, R_7 = 2\text{-ethylhexyl}, R_8 = \text{diphenylamino}$



- polymer 43 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = \text{t-butyl}$
- polymer 44 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = \text{H}$
- polymer 45 $R_5 = R_6 = \text{n-hexyl}, R_7 = (4\text{-carbazole})\text{phenyl}$
- 5 polymer 46 $R_5 = \text{n-hexyloxy}, R_6 = \text{H}, R_7 = \text{diphenylamino}$
- polymer 47 $R_5 = R_6 = \text{n-hexyl}, R_7 = \text{trifluoromethyl}$

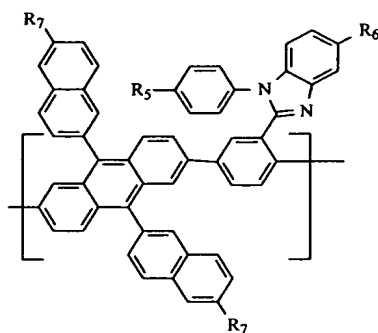


- polymer 48 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = \text{t-butyl}$
- 10 polymer 49 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = \text{H}$
- polymer 50 $R_5 = R_6 = \text{n-hexyl}, R_7 = (4\text{-carbazole})\text{phenyl}$
- polymer 51 $R_5 = R_7 = \text{n-hexyloxy}, R_6 = \text{diphenylamino}$
- polymer 52 $R_5 = \text{H}, R_6 = \text{n-hexyl}, R_7 = \text{trifluoromethyl}$



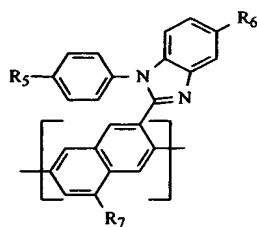
15

- polymer 53 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = \text{t-butyl}$
polymer 54 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = \text{methoxycarbonyl}$
polymer 55 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = R_7 = \text{H}$
polymer 56 $R_5 = R_6 = \text{n-hexyl}, R_7 = (4\text{-carbazole})\text{phenyl}$
5 polymer 57 $R_5 = \text{n-hexyloxy}, R_6 = \text{H}, R_7 = \text{diphenylamino}$
polymer 58 $R_5 = R_6 = \text{n-hexyl}, R_7 = \text{trifluoromethyl}$

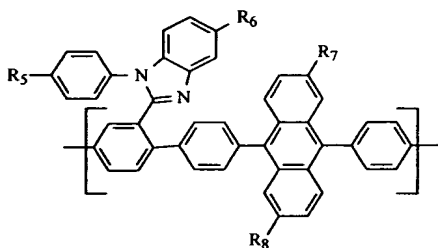


- polymer 59 $R_5 = R_6 = R_7 = \text{n-hexyl}$
10 polymer 60 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = 2\text{-ethylhexyloxy}$
polymer 61 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = \text{n-hexyl}, R_7 = \text{t-butyl}$
polymer 62 $R_5 = R_6 = \text{n-hexyl}, R_7 = (4\text{-carbazole})\text{phenyl}$
polymer 63 $R_5 = R_7 = \text{n-hexyloxy}, R_6 = \text{diphenylamino}$

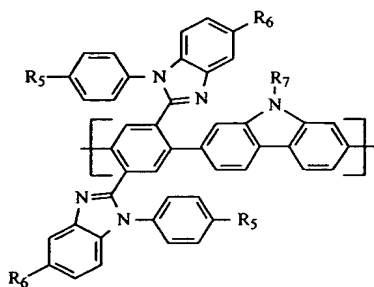
15



- polymer 64 $R_5 = R_6 = R_7 = \text{n-hexyl}$
polymer 65 $R_5 = \text{n-hexyl}, R_6 = \text{H}, R_7 = 2\text{-ethylhexyloxy}$
polymer 66 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}, R_6 = \text{n-hexyl}, R_7 = \text{H}$
20 polymer 67 $R_5 = R_6 = \text{n-hexyl}, R_7 = (4\text{-carbazole})\text{phenyl}$
polymer 68 $R_5 = R_7 = \text{n-hexyloxy}, R_6 = \text{diphenylamino}$

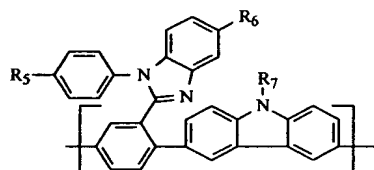


- polymer 69 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = R_8 = 2\text{-ethylhexyloxy}$
 polymer 70 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = R_8 = \text{t-butyl}$
 polymer 71 $R_5 = R_7 = R_8 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 5 polymer 72 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = 2\text{-ethylhexyl}$, $R_8 = \text{diphenylamino}$
 polymer 73 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = R_8 = 3,7\text{-dimethyloctyl}$



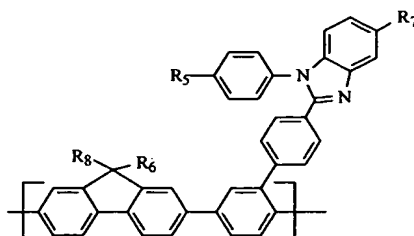
- polymer 74 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
 10 polymer 75 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = 2\text{-ethylhexyl}$
 polymer 76 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 polymer 77 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$
 polymer 78 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{t-butyl}$
 polymer 79 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{trifluoromethyl}$

15



- polymer 80 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
 polymer 81 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 polymer 82 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

polymer 83 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{t-butyl}$



polymer 84 $R_5 = R_6 = R_7 = R_8 = \text{n-hexyl}$

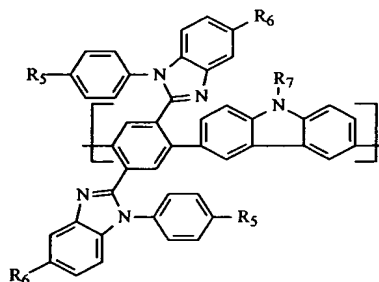
5 polymer 85 $R_5 = R_7 = \text{H}$, $R_6 = R_8 = 3,7\text{-dimethyloctyl}$

polymer 86 $R_5 = R_7 = \text{H}$, $R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_8 = \text{n-hexyl}$

polymer 87 $R_5 = 4\text{-(N-carbazole)phenyl}$, $R_6 = R_8 = \text{n-decyl}$, $R_7 = \text{H}$

polymer 88 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = \text{n-hexyl}$, $R_8 = \text{n-octyl}$

10

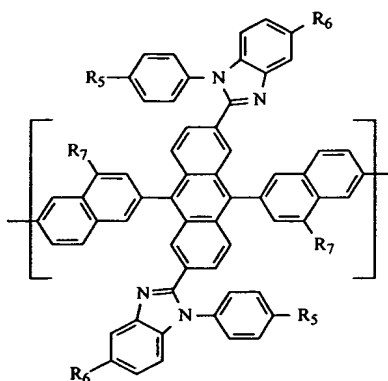


polymer 89 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$

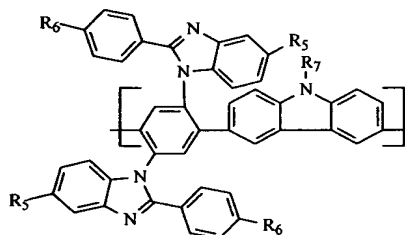
polymer 90 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = 2\text{-ethylhexyl}$

polymer 91 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

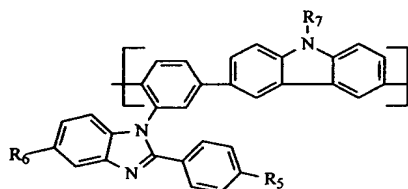
15 polymer 92 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$



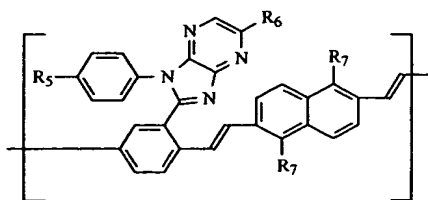
- polymer 93 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
polymer 94 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
polymer 95 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$
5 polymer 96 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$



- polymer 97 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
polymer 98 $R_5 = \text{H}$, $R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$
10 polymer 99 $R_5 = \text{H}$, $R_6 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$
polymer 100 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$



- polymer 101 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
15 polymer 102 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = 2\text{-ethylhexyl}$
polymer 103 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

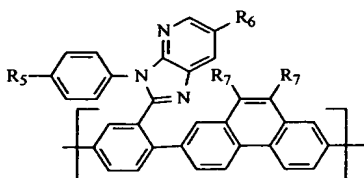


polymer 104 $R_5 = R_6 = R_7 = \text{n-hexyl}$

polymer 105 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

polymer 106 $R_5 = \text{H}$, $R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

5 polymer 107 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$

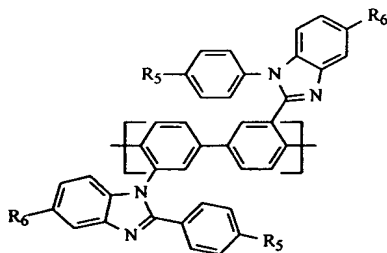


polymer 108 $R_5 = R_6 = 3,7\text{-dimethyloctyl}$, $R_7 = \text{H}$

polymer 109 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

10 polymer 110 $R_5 = R_6 = \text{n-hexyloxy}$, $R_7 = 2\text{-ethylhexyloxy}$

polymer 111 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{t-butyl}$



polymer 112 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$

15 polymer 113 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$

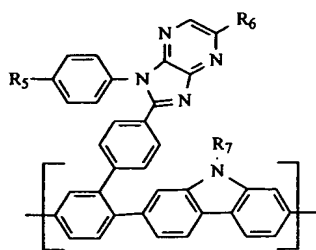
polymer 114 $R_5 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{n-hexyl}$

polymer 115 $R_5 = R_6 = \text{n-hexyl}$

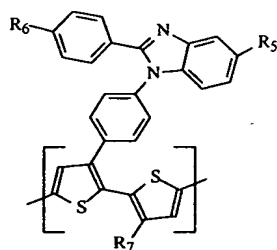
polymer 116 $R_5 = \text{H}$, $R_6 = \text{n-hexyloxy}$

polymer 117 $R_5 = \text{n-hexyl}$, $R_6 = 4\text{-(t-butylphenyl)}$

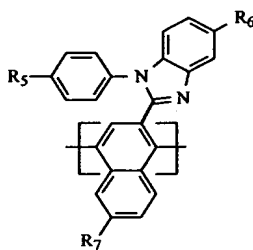
20 polymer 118 $R_5 = R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$



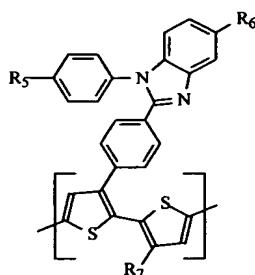
- polymer 119 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
 polymer 120 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = 4\text{-(t-butylphenyl)}$
 polymer 121 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 5 polymer 122 $R_5 = \text{H}$, $R_6 = \text{n-hexyl}$, $R_7 = 3,7\text{-dimethyloctyl}$



- polymer 123 $R_5 = \text{n-hexyl}$, $R_6 = R_7 = \text{H}$
 polymer 124 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 10 polymer 125 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$
 polymer 126 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = \text{H}$



- polymer 127 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$
 15 polymer 128 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$
 polymer 129 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$
 polymer 130 $R_5 = \text{H}$, $R_6 = \text{n-hexyl}$, $R_7 = 3,7\text{-dimethyloctyl}$

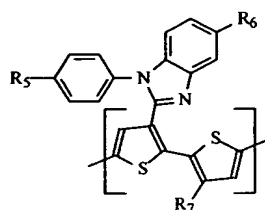


polymer 131 $R_5 = \text{n-hexyl}$, $R_6 = R_7 = \text{H}$

polymer 132 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

polymer 133 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

5 polymer 134 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = \text{H}$

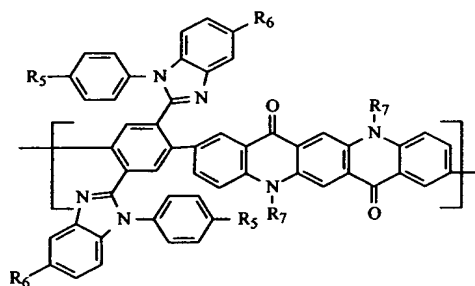


polymer 135 $R_5 = \text{n-hexyl}$, $R_6 = R_7 = \text{H}$

polymer 136 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

10 polymer 137 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

polymer 138 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = \text{H}$



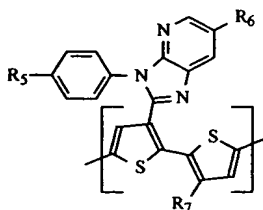
polymer 139 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$

15 polymer 140 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

polymer 141 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

polymer 142 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{t-butyl}$

polymer 143 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{trifluoromethyl}$



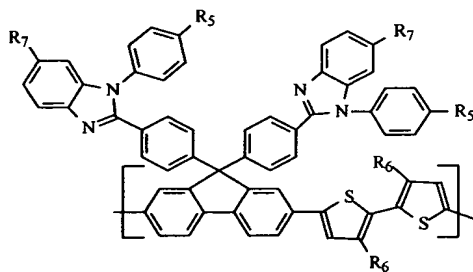
polymer 144 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$

polymer 145 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

5 polymer 146 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

polymer 147 $R_5 = \text{n-hexyloxy}$, $R_6 = R_7 = 2\text{-ethylhexyl}$

polymer 148 $R_5 = R_6 = R_7 = \text{n-hexyl}$



10 polymer 149 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-t-butyl})\text{phenyl}$

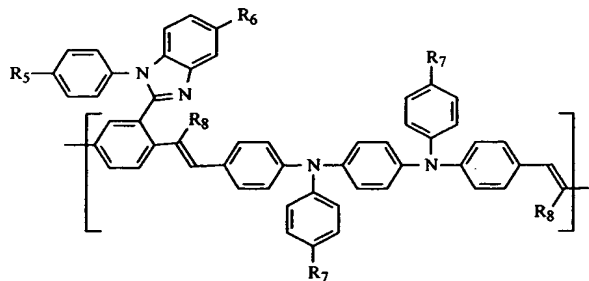
polymer 150 $R_5 = R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$

polymer 151 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = 2\text{-ethylhexyl}$

polymer 152 $R_5 = \text{n-hexyloxy}$, $R_6 = 2\text{-ethylhexyl}$, $R_7 = \text{t-butyl}$

polymer 153 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{trifluoromethyl}$

15

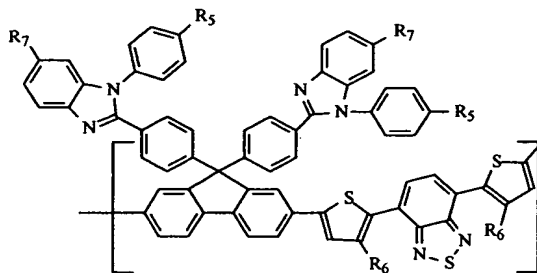


polymer 154 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{t-butyl}$, $R_8 = \text{H}$

polymer 155 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{n-hexyl}$, $R_7 = 4\text{-t-butylphenyl}$, $R_8 = \text{CN}$

polymer 156 $R_5 = R_6 = n\text{-hexyloxy}$, $R_7 = t\text{-butyl}$, $R_8 = \text{phenyl}$

polymer 157 $R_5 = n\text{-hexyl}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$, $R_8 = CN$

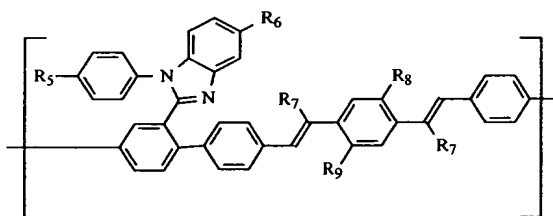


5 polymer 158 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = t\text{-butyl}$

polymer 159 $R_5 = R_6 = 2\text{-ethylhexyl}$, $R_7 = 4\text{-}t\text{-butylphenyl}$

polymer 160 $R_5 = n\text{-hexyloxy}$, $R_6 = n\text{-hexyl}$, $R_7 = H$

polymer 161 $R_5 = n\text{-hexyl}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$



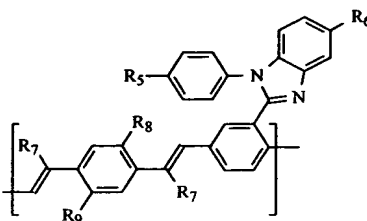
10

polymer 162 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{phenyl}$, $R_8 = R_9 = 2\text{-ethylhexyl}$

polymer 163 $R_5 = 2\text{-ethylhexyl}$, $R_6 = n\text{-hexyl}$, $R_7 = R_8 = H$, $R_9 = 4\text{-}t\text{-butylphenyl}$

polymer 164 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = H$, $R_8 = \text{methoxy}$, $R_9 = 3,7\text{-dimethyloctyloxy}$

15 polymer 165 $R_5 = n\text{-hexyl}$, $R_6 = R_8 = 3,7\text{-dimethyloctyl}$, $R_7 = H$, $R_9 = (4\text{-diphenylamino})\text{phenyl}$



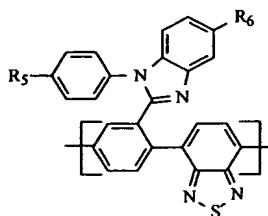
polymer 166 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{phenyl}$, $R_8 = R_9 = 2\text{-ethylhexyl}$

polymer 167 $R_5 = 2\text{-ethylhexyl}$, $R_6 = n\text{-hexyl}$, $R_7 = R_8 = H$, $R_9 = 4\text{-}t\text{-butylphenyl}$

polymer 168 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = H$, $R_8 = \text{methoxy}$, $R_9 = 3,7\text{-dimethyloctyloxy}$

polymer 169 $R_5 = n\text{-hexyl}$, $R_6 = R_8 = 3,7\text{-dimethyloctyl}$, $R_7 = H$, $R_9 = (4\text{-diphenylamino})\text{phenyl}$

polymer 170 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{phenyl}$, $R_8 = R_9 = 2\text{-ethylhexyloxy}$



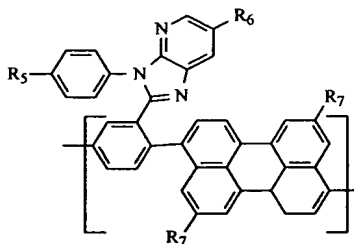
polymer 171 $R_5 = R_6 = n\text{-hexyl}$

polymer 172 $R_5 = n\text{-hexyloxy}$, $R_6 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$

polymer 173 $R_5 = n\text{-hexyl}$, $R_6 = 2\text{-ethylhexyl}$

polymer 174 $R_5 = H$, $R_6 = n\text{-hexyloxy}$

polymer 175 $R_5 = \text{trifluoromethyl}$, $R_6 = 3,7\text{-dimethyloctyl}$

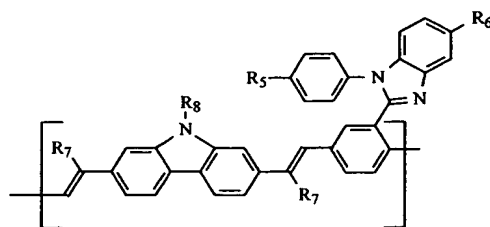


polymer 176 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = t\text{-butyl}$

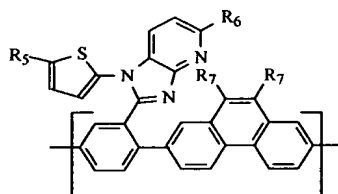
polymer 177 $R_5 = R_6 = 2\text{-ethylhexyl}$, $R_7 = 4\text{-}t\text{-butylphenyl}$

polymer 178 $R_5 = n\text{-hexyloxy}$, $R_6 = n\text{-hexyl}$, $R_7 = H$

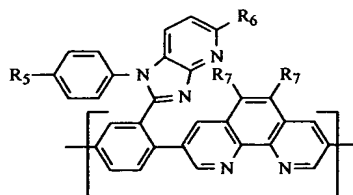
polymer 179 $R_5 = n\text{-hexyl}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$



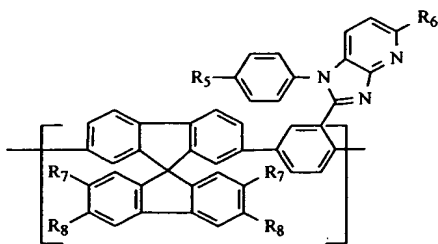
- polymer 180 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{phenyl}$, $R_8 = 2\text{-ethylhexyl}$
 polymer 181 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{n-hexyl}$, $R_7 = \text{H}$, $R_8 = 4\text{-t-butylphenyl}$
 polymer 182 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = \text{CN}$, $R_8 = 3,7\text{-dimethyloctyloxy}$
 5 polymer 183 $R_5 = \text{n-hexyloxy}$, $R_6 = R_8 = 3,7\text{-dimethyloctyl}$, $R_7 = \text{H}$



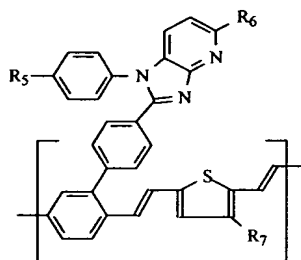
- polymer 184 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{t-butyl}$
 polymer 185 $R_5 = R_6 = 2\text{-ethylhexyl}$, $R_7 = 4\text{-t-butylphenyl}$
 10 polymer 186 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$



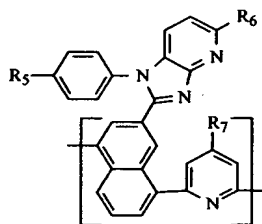
- polymer 187 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{t-butyl}$
 polymer 188 $R_5 = R_6 = 2\text{-ethylhexyl}$, $R_7 = 4\text{-t-butylphenyl}$
 15 polymer 189 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{n-octyl}$
 polymer 190 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$
 polymer 191 $R_5 = R_7 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $(4\text{-diphenylamino})\text{phenyl}$



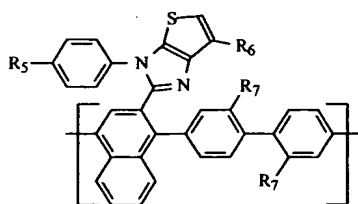
- polymer 192 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{H}$, $R_8 = \text{2-ethylhexyloxy}$
 polymer 193 $R_5 = \text{2-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = R_8 = \text{2-ethylhexyloxy}$
 polymer 194 $R_5 = R_6 = \text{n-hexyloxy}$, $R_7 = \text{H}$, $R_8 = \text{3,7-dimethyloctyloxy}$
 5 polymer 195 $R_5 = (\text{4-diphenylamino})\text{phenyl}$, $R_6 = R_7 = R_8 = \text{3,7-dimethyloctyl}$



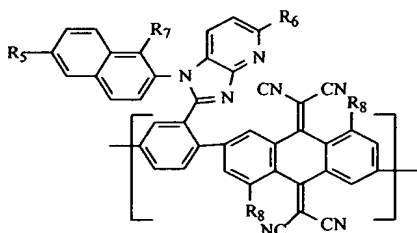
- polymer 196 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{t-butyl}$
 polymer 197 $R_5 = R_6 = \text{2-ethylhexyl}$, $R_7 = \text{4-t-butylphenyl}$
 10 polymer 198 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{n-octyl}$
 polymer 199 $R_5 = \text{n-hexyl}$, $R_6 = \text{H}$, $R_7 = (\text{4-diphenylamino})\text{phenyl}$



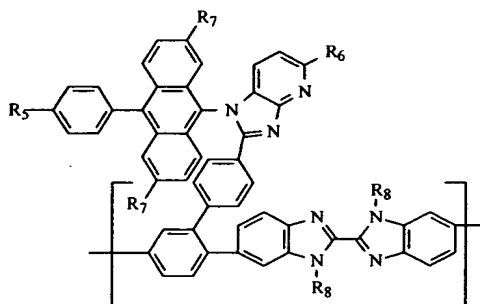
- polymer 200 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{dimethylamino}$
 15 polymer 201 $R_5 = R_6 = \text{2-ethylhexyl}$, $R_7 = \text{4-t-butylphenyl}$
 polymer 202 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{n-octyl}$
 polymer 203 $R_5 = \text{H}$, $R_6 = \text{n-hexyl}$, $R_7 = (\text{4-diphenylamino})\text{phenyl}$



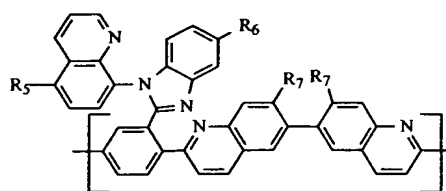
- polymer 204 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{dimethylamino}$
 polymer 205 $R_5 = R_7 = \text{2-ethylhexyloxy}$, $R_6 = \text{2-ethylhexyl}$
 polymer 206 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{n-octyl}$
 5 polymer 207 $R_5 = \text{H}$, $R_6 = R_7 = \text{(4-diphenylamino)phenyl}$



- polymer 208 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{(4-t-butyl)phenyl}$, $R_8 = \text{H}$
 polymer 209 $R_5 = R_7 = \text{4-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$, $R_8 = \text{hexyloxy}$
 10 polymer 210 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$, $R_8 = \text{2-ethylhexyl}$
 polymer 211 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{2-ethylhexyl}$, $R_7 = \text{t-butyl}$, $R_8 = \text{H}$



- 15 polymer 212 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{2-ethylhexyloxy}$, $R_8 = \text{phenyl}$
 polymer 213 $R_5 = R_7 = \text{4-(bis(4-methylphenyl)amino)phenyl}$, $R_6 = \text{H}$, $R_8 = \text{hexyl}$
 polymer 214 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$, $R_8 = \text{2-ethylhexyl}$
 polymer 215 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{2-ethylhexyl}$, $R_7 = \text{t-butyl}$, $R_8 = \text{methyl}$

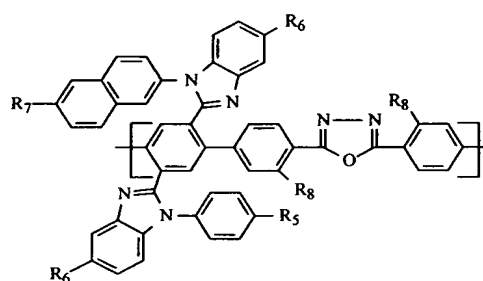


polymer 216 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{dimethylamino}$

polymer 217 $R_5 = R_6 = \text{2-ethylhexyloxy}$, $R_7 = \text{4-t-butylphenyl}$

polymer 218 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{n-octyl}$

5 polymer 219 $R_5 = \text{H}$, $R_6 = R_7 = (\text{4-diphenylamino})\text{phenyl}$



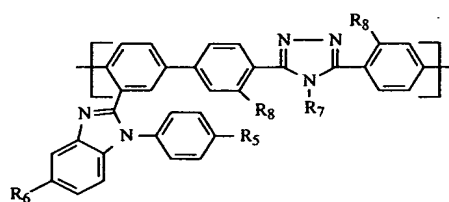
polymer 220 $R_5 = R_6 = R_7 = \text{n-hexyl}$, $R_8 = (\text{4-t-butyl})\text{phenyl}$

polymer 221 $R_5 = R_6 = \text{hexyloxy}$, $R_7 = \text{4-(bis(4-methylphenyl)amino)phenyl}$, R_8

10 = H

polymer 222 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$, $R_8 = \text{2-ethylhexyl}$

polymer 223 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{2-ethylhexyl}$, $R_7 = \text{H}$, $R_8 = \text{di-tolylamino}$

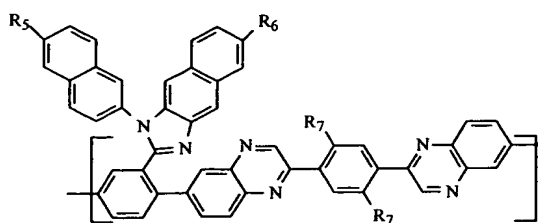


15 polymer 224 $R_5 = R_6 = R_7 = \text{n-hexyl}$, $R_8 = (\text{4-t-butyl})\text{phenyl}$

polymer 225 $R_5 = R_6 = \text{hexyloxy}$, $R_7 = \text{4-(bis(4-methylphenyl)amino)phenyl}$, R_8
= H

polymer 226 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{2-ethylhexyl}$, $R_7 = \text{4-(di-tolylamino)phenyl}$,
 $R_8 = \text{H}$

20

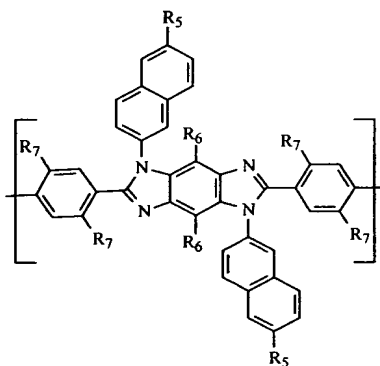


polymer 227 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{dimethylamino}$

polymer 228 $R_5 = R_6 = R_7 = 2\text{-ethylhexyloxy}$, 4-t-butylphenyl

polymer 229 $R_5 = n\text{-hexyloxy}$, $R_6 = H$, $R_7 = n\text{-octyl}$

5 polymer 230 $R_5 = H$, $R_6 = R_7 = (4\text{-ditolylamino})\text{phenyl}$

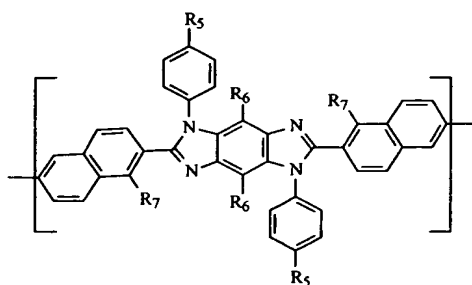


polymer 231 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{hexyloxy}$

polymer 232 $R_5 = 2\text{-ethylhexyl}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

10 polymer 233 $R_5 = n\text{-hexyloxy}$, $R_6 = \text{octyl}$, $R_7 = H$

polymer 234 $R_5 = n\text{-hexyl}$, $R_6 = H$, $R_7 = 4\text{-octylphenyl}$

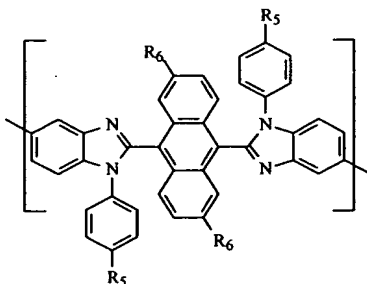


polymer 235 $R_5 = R_6 = R_7 = n\text{-hexyl}$

15 polymer 236 $R_5 = 2\text{-ethylhexyl}$, $R_6 = R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 237 $R_5 = n\text{-hexyloxy}$, $R_6 = \text{octyl}$, $R_7 = H$

polymer 238 $R_5 = \text{di-tolylamino}$, $R_6 = \text{H}$, $R_7 = 4\text{-octyl}$

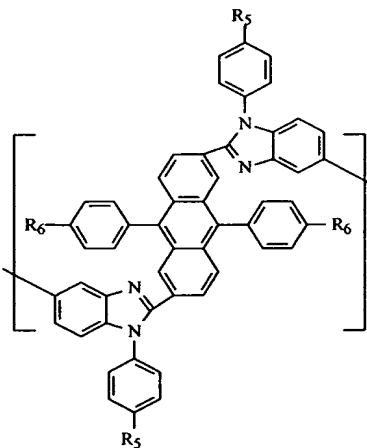


polymer 239 $R_5 = R_6 = \text{n-hexyl}$

5 polymer 240 $R_5 = 2\text{-ethylhexyl}$, $R_6 = (4\text{-diphenylamino})\text{phenyl}$

polymer 241 $R_5 = R_6 = \text{n-hexyloxy}$

polymer 242 $R_5 = \text{di-tolylamino}$, $R_6 = \text{octyloxy}$



10 polymer 243 $R_5 = R_6 = \text{n-hexyl}$

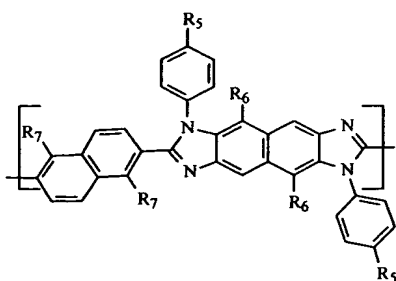
polymer 244 $R_5 = R_6 = \text{n-octyloxy}$

polymer 245 $R_5 = 2\text{-ethylhexyl}$, $R_6 = (4\text{-diphenylamino})\text{phenyl}$

polymer 246 $R_5 = \text{n-hexyloxy}$, $R_6 = \text{octyl}$

polymer 247 $R_5 = 4\text{-octylphenyl}$, $R_6 = \text{trifluoromethyl}$

15

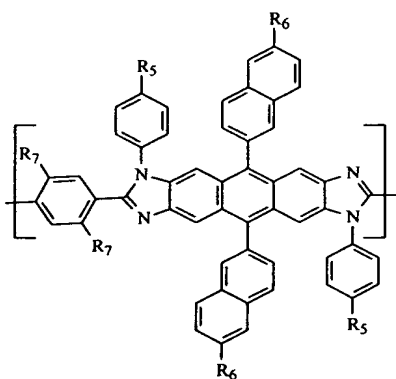


polymer 248 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{hexyloxy}$

polymer 249 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 250 $R_5 = n\text{-hexyloxy}$, $R_6 = \text{octyl}$, $R_7 = \text{H}$

5 polymer 251 $R_5 = \text{H}$, $R_6 = \text{octyloxy}$, $R_7 = 4\text{-octylphenyl}$



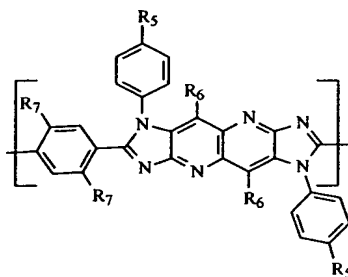
polymer 252 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{hexyloxy}$

polymer 253 $R_5 = R_6 = R_7 = \text{hexyloxy}$

10 polymer 254 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 255 $R_5 = \text{H}$, $R_6 = 3,7\text{-dimethyloctyl}$, $R_7 = \text{hexyloxy}$

polymer 256 $R_5 = \text{ditolylamino}$, $R_6 = 4\text{-octylphenyl}$, $R_7 = \text{H}$

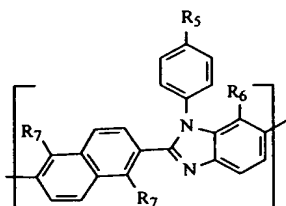


15 polymer 257 $R_5 = R_6 = n\text{-hexyl}$, $R_7 = \text{hexyloxy}$

polymer 258 $R_5 = R_6 = R_7 = \text{hexyloxy}$

polymer 259 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 260 $R_5 = \text{H}$, $R_6 = \text{hexyloxy}$, $R_7 = 3,7\text{-dimethyloctyl}$



5

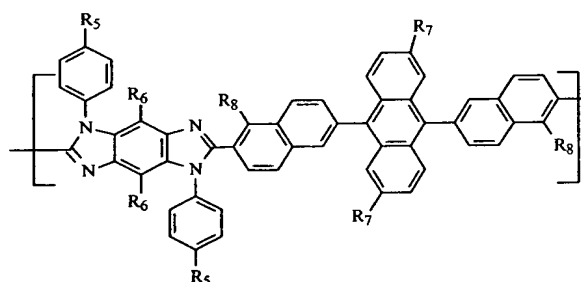
polymer 261 $R_5 = R_6 = \text{n-hexyl}$, $R_7 = \text{hexyloxy}$

polymer 262 $R_5 = R_6 = R_7 = \text{hexyloxy}$

polymer 263 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 264 $R_5 = \text{H}$, $R_6 = \text{hexyloxy}$, $R_7 = 3,7\text{-dimethyloctyl}$

10

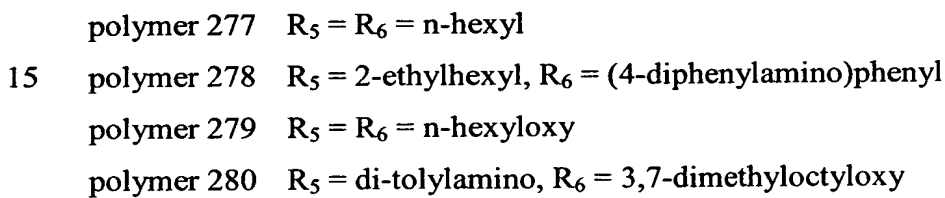
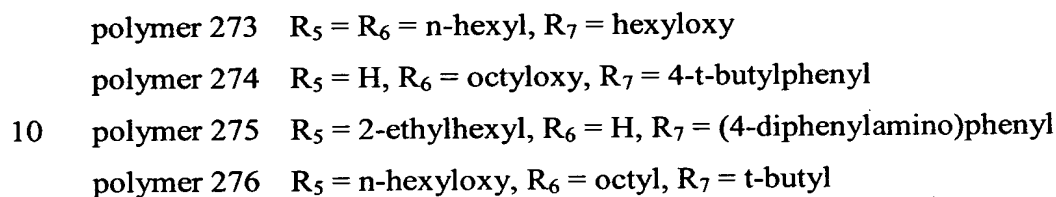
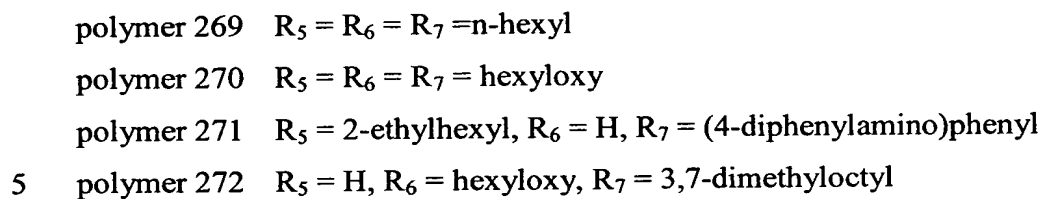


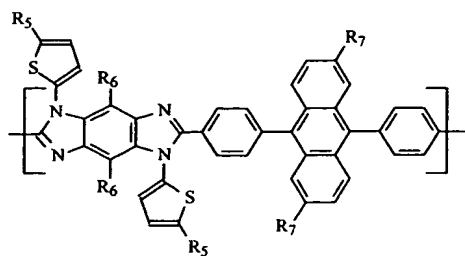
polymer 265 $R_5 = R_6 = R_7 = \text{n-hexyl}$, $R_8 = \text{t-butyl}$

polymer 266 $R_5 = R_6 = \text{hexyloxy}$, $R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, $R_8 = \text{H}$

15 polymer 267 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$, $R_8 = 2\text{-ethylhexyl}$

polymer 268 $R_5 = \text{H}$, $R_6 = 2\text{-ethylhexyl}$, $R_7 = 3,7\text{-dimethyloctyl}$, $R_8 = \text{di-tolylamino}$



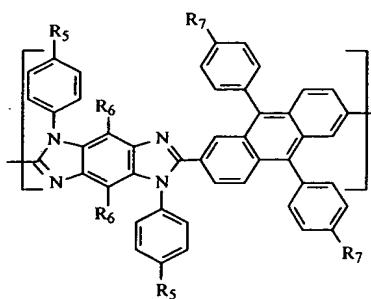


polymer 281 $R_5 = R_6 = R_7 = \text{n-hexyl}$

polymer 282 $R_5 = 2\text{-ethylhexyl}, R_6 = \text{H}, R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 283 $R_5 = \text{H}, R_6 = \text{hexyloxy}, R_7 = 3,7\text{-dimethyloctyl}$

5

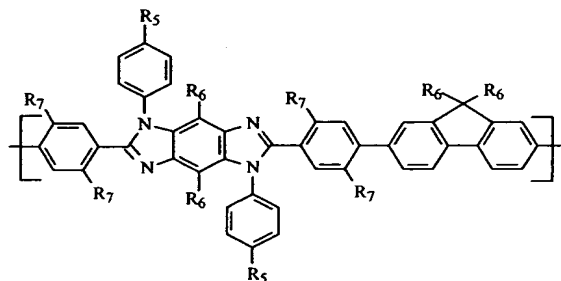


polymer 284 $R_5 = R_6 = R_7 = \text{n-hexyl}$

polymer 285 $R_5 = R_6 = R_7 = \text{hexyloxy}$

polymer 286 $R_5 = 2\text{-ethylhexyl}, R_6 = \text{H}, R_7 = (4\text{-diphenylamino})\text{phenyl}$

10 polymer 287 $R_5 = \text{H}, R_6 = \text{hexyloxy}, R_7 = 3,7\text{-dimethyloctyloxy}$

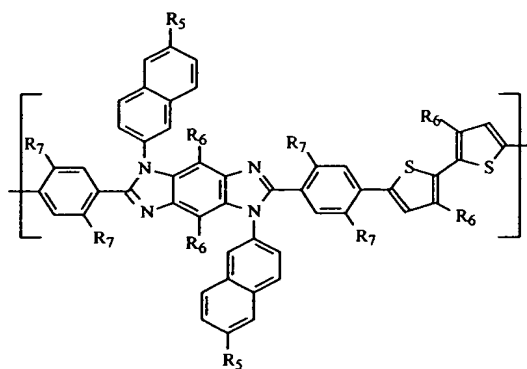


polymer 288 $R_5 = R_6 = \text{n-hexyl}, R_7 = \text{hexyloxy}$

polymer 289 $R_5 = \text{H}, R_6 = \text{tolyl}, R_7 = 4\text{-octylphenyl}$

15 polymer 290 $R_5 = 2\text{-ethylhexyl}, R_6 = (4\text{-diphenylamino})\text{phenyl}, R_7 = \text{H}$

polymer 291 $R_5 = \text{n-hexyloxy}, R_6 = \text{octyl}, R_7 = \text{t-butyl}$

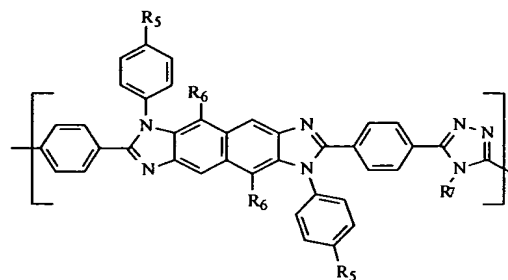


polymer 292 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 293 $R_5 = 2\text{-ethylhexyloxy}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 294 $R_5 = R_7 = 3,7\text{-dimethyloctyl}$, $R_6 = H$

5

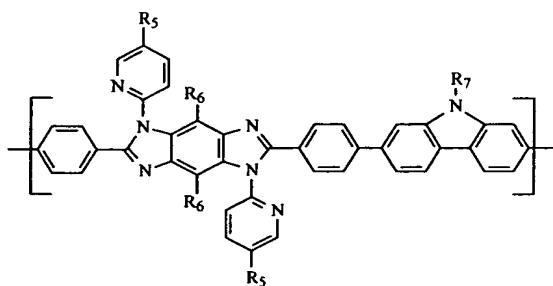


polymer 295 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 296 $R_5 = 2\text{-ethylhexyl}$, $R_6 = H$, $R_7 = (4\text{-diphenylamino})\text{phenyl}$

polymer 297 $R_5 = H$, $R_6 = \text{hexyloxy}$, $R_7 = 3,7\text{-dimethyloctyloxy}$

10

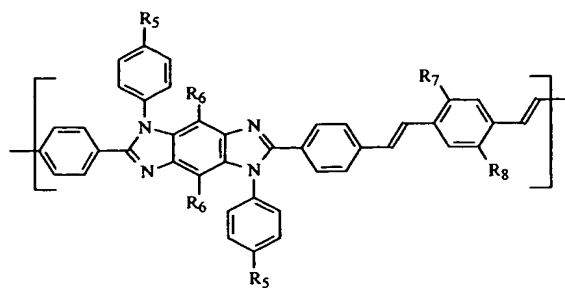


polymer 298 $R_5 = R_6 = R_7 = n\text{-hexyl}$,

polymer 299 $R_5 = H$, $R_6 = \text{octyl}$, $R_7 = 4\text{-octylphenyl}$

polymer 300 $R_5 = 2\text{-ethylhexyl}$, $R_6 = H$, $R_7 = \text{ditolylaminophenyl}$

15 polymer 301 $R_5 = n\text{-hexyloxy}$, $R_6 = R_7 = \text{octyl}$



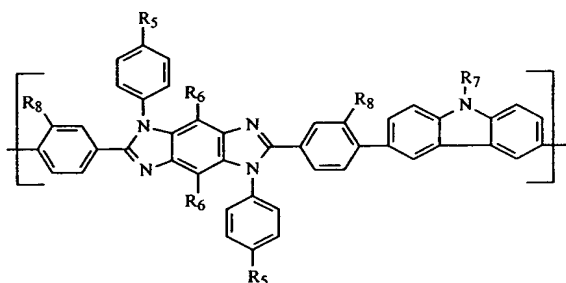
polymer 302 $R_5 = R_6 = R_7 = R_8 = \text{n-hexyl}$

polymer 303 $R_5 = R_6 = \text{hexyloxy}$, $R_7 = 4\text{-(bis(4-methylphenyl)amino)phenyl}$, R_8

5 = H

polymer 304 $R_5 = R_6 = R_7 = \text{n-hexyloxy}$, $R_8 = \text{methoxy}$

polymer 305 $R_5 = \text{H}$, $R_6 = 2\text{-ethylhexyl}$, $R_7 = 3,7\text{-dimethyloctyl}$, $R_8 = \text{trifluoromethyl}$

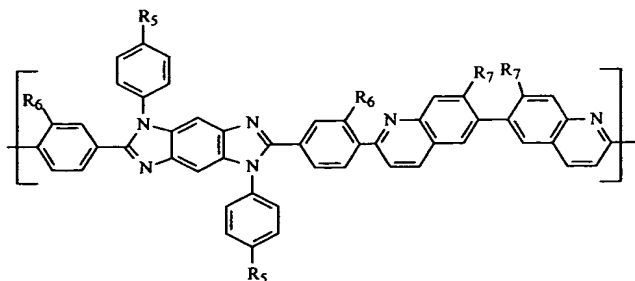


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polymer 306 $R_5 = R_6 = R_7 = \text{n-hexyl}$, $R_8 = \text{H}$

polymer 307 $R_5 = R_6 = R_8 = \text{hexyloxy}$, $R_7 = 4\text{-(t-butylphenyl)}$

polymer 308 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = 3,7\text{-dimethyloctyl}$, $R_8 = \text{di-tolylamino}$



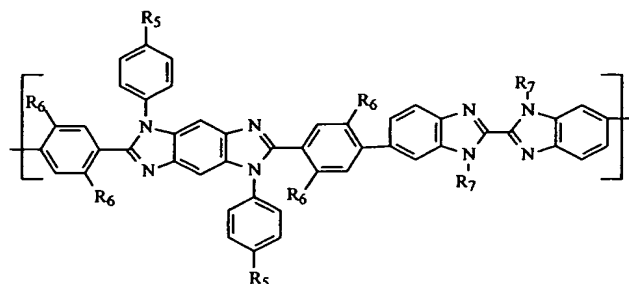
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polymer 309 $R_5 = R_6 = R_7 = \text{n-hexyl}$

polymer 310 $R_5 = R_6 = \text{octyloxy}$, $R_7 = 4\text{-octylphenyl}$

polymer 311 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = \text{ditolylaminophenyl}$

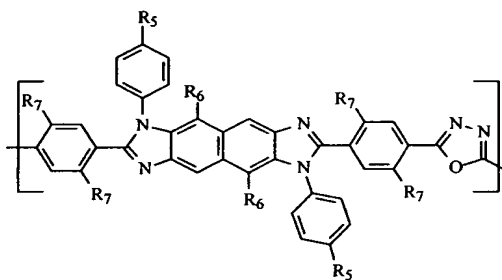
polymer 312 $R_5 = n\text{-hexyloxy}$, $R_6 = R_7 = \text{octyl}$



5 polymer 313 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 314 $R_5 = R_6 = \text{octyloxy}$, $R_7 = 4\text{-octylphenyl}$

polymer 315 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = \text{octyl}$

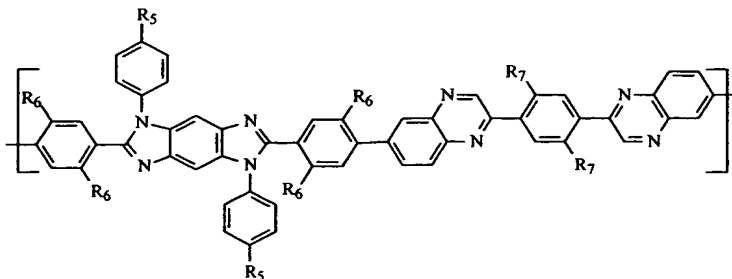


10 polymer 316 $R_5 = R_6 = R_7 = n\text{-hexyl}$,

polymer 317 $R_5 = \text{hexyl}$, $R_6 = \text{octyloxy}$, $R_7 = 4\text{-t-butylphenyl}$

polymer 318 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = \text{octyloxy}$

polymer 319 $R_5 = R_6 = \text{H}$, $R_7 = \text{octyloxy}$



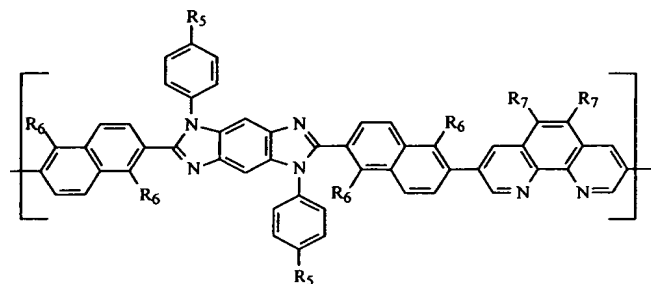
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polymer 320 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 321 $R_5 = R_6 = R_7 = \text{octyloxy}$

polymer 322 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{ditolylaminophenyl}$, $R_7 = \text{H}$

polymer 323 $R_5 = n\text{-hexyloxy}$, $R_6 = \text{H}$, $R_7 = \text{octyl}$

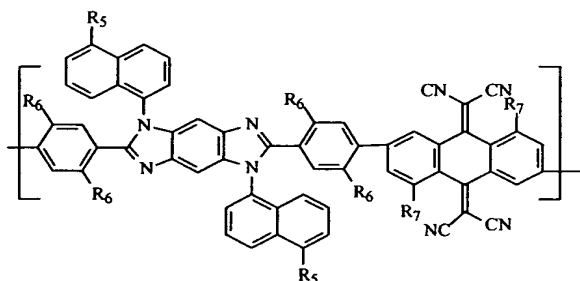


5 polymer 324 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{hexyloxy}$, $R_7 = t\text{-butyl}$

polymer 325 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 326 $R_5 = R_6 = R_7 = \text{octyloxy}$

polymer 327 $R_5 = \text{methyl}$, $R_6 = 3,7\text{-dimethyloctyl}$, $R_7 = \text{ditolylaminophenyl}$



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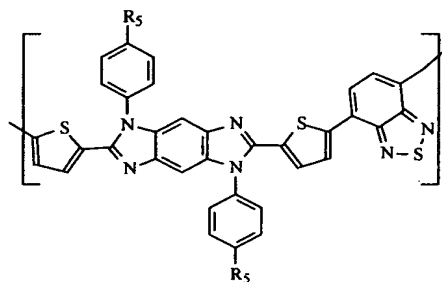
polymer 328 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 329 $R_5 = R_6 = R_7 = \text{hexyloxy}$

polymer 330 $R_5 = 2\text{-ethylhexyl}$, $R_6 = (4\text{-diphenylamino})\text{phenyl}$, $R_7 = \text{H}$

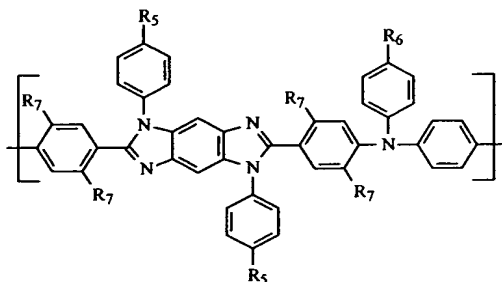
polymer 331 $R_5 = R_7 = \text{H}$, $R_6 = 3,7\text{-dimethyloctyloxy}$

15



polymer 332 $R_5 = n\text{-hexyl}$

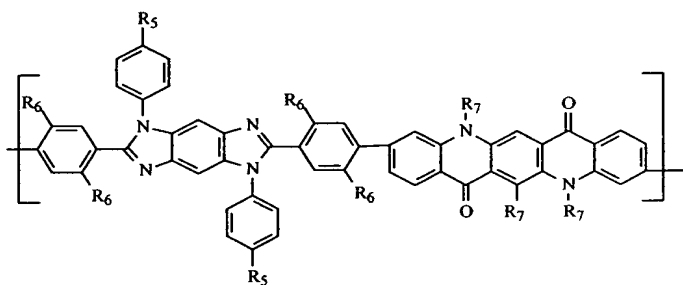
- polymer 333 $R_5 = 3,7\text{-dimethyloctyloxy}$
 polymer 334 $R_5 = \text{bis}(4\text{-octylphenylamino})$
 polymer 335 $R_5 = \text{H}, R_6 = \text{hexyloxy}$



5

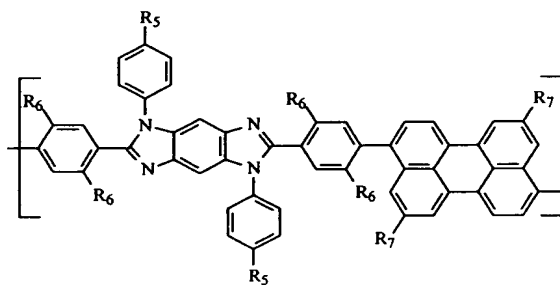
- polymer 336 $R_5 = R_6 = R_7 = n\text{-hexyl}$
 polymer 337 $R_5 = R_6 = R_7 = \text{hexyloxy}$
 polymer 338 $R_5 = 2\text{-ethylhexyl}, R_6 = (4\text{-di-tolylamino}), R_7 = \text{H}$
 polymer 339 $R_5 = \text{H}, R_6 = \text{hexyloxy}, R_7 = 3,7\text{-dimethyloctyloxy}$

10



- polymer 340 $R_5 = R_6 = R_7 = n\text{-hexyl}$
 polymer 341 $R_5 = 2\text{-ethylhexyl}, R_6 = \text{H}, R_7 = 4\text{-octylphenyl}$
 polymer 342 $R_5 = \text{H}, R_6 = \text{hexyloxy}, R_7 = 3,7\text{-dimethyloctyl}$

15



- polymer 343 $R_5 = R_6 = R_7 = n\text{-hexyl}$

polymer 344 $R_5 = R_6 = R_7 = \text{hexyloxy}$

polymer 345 $R_5 = 2\text{-ethylhexyl}$, $R_6 = \text{H}$, $R_7 = 4\text{-octylphenyl}$

polymer 346 $R_5 = \text{H}$, $R_6 = (4\text{-octylphenyl})\text{amino}$, $R_7 = 3,7\text{-dimethyloctyloxy}$

5 The specific molecular structures can be the combination of any of the above drawn structures.

The conjugated polymers comprising azole structure (I) can be synthesized using known methods. The polymerization method and the molecular weights of the resulting polymers used in the present invention are not necessary to be particularly restricted. The molecular weights of the polymers are at least
10 1000, and preferably at least 2000. The polymers may be prepared by condensation polymerizations, such as coupling reactions including Pd-catalyzed Suzuki coupling, Stille coupling or Heck coupling, or Ni-mediated Yamamoto coupling, or by other condensation methods such as Wittig reaction, or Horner-Emmons reaction, or Knoevenagel reaction, or dehalogenation of dibenzyl
15 halides. Preferably the polymers are prepared by Suzuki coupling reaction.

Suzuki coupling reaction was first reported by Suzuki et al on the coupling of aromatic boronic acid derivatives with aromatic halides (Suzuki, A. et al *Synthetic Comm.* 1981, 11(7), 513). Since then, this reaction has been widely used to prepared polymers for various applications (Ranger, M. et al
20 *Macromolecules* 1997, 30, 7686). The reaction involves the use of a palladium-based catalyst such as a soluble Pd compound either in the state of Pd (II) or Pd (0), a base such as an aqueous inorganic alkaline carbonate or bicarbonate, and a solvent for the reactants and/or product. The preferred Pd catalyst is a Pd (0) complex such as $\text{Pd}(\text{PPh}_3)_4$ or a Pd (II) salt such as $\text{Pd}(\text{PPh}_3)_2\text{Cl}_2$ or $\text{Pd}(\text{OAc})_2$
25 with a tertiary phosphine ligand, and used in the range of 0.01-10 mol% based on the functional groups of the reactants. Polar solvents such as THF and non-polar solvents toluene can be used however, the non-polar solvent is believed to slow down the reaction. Modified processes were reported to prepare conjugated polymers for EL devices from the Suzuki coupling of aromatic halides and
30 aromatic boron derivatives (Inbasekaran, M. et al U.S. Patent No. 5,777,070 (1998); Towns, C. R. et al. PCT WO00/53656, 2000). A variation of the Suzuki

coupling reaction replaces the aromatic halide with an aromatic trifluoromethanesulfonate (triflate) (Ritter, K. Synthesis, 1993, 735). Aromatic triflates are readily prepared from the corresponding phenol derivatives. The advantages of using aromatic triflates are that the phenol derivatives are easily
5 accessible and can be protected/deprotected during complex synthesis. For example, aromatic halides normally would react under various coupling conditions to generate unwanted by-product and lead to much more complicated synthetic schemes. However, phenol derivatives can be easily protected by various protecting groups that would not interfere with functional group
10 transformation and be deprotected to generate back the phenol group which then can be converted to triflates. The diboron derivatives can be prepared from the corresponding dihalide or ditriflate.

The synthetic schemes of the polymers according to the present invention are illustrated in Schemes 1-4.

15 The process of the invention provides conjugated polymers particularly useful for an optical device. The optical device may include a luminescent device such as an EL device in which the polymers of the present invention is deposited between a cathode and an anode. The polymers can be deposited as thin film by vapor deposition method or thermal transfer, or from a
20 solution by spin-coating, spray-coating, dip-coating, roller-coating, or ink jet delivery. The thin film may be supported by substrate directly, preferably a transparent substrate, or supported by the substrate indirectly where there is one or more inter layers between the substrate and thin film. The thin film can be used as emitting layer or charge carrier transporting layer, especially electron transporting
25 layer.

General EL device architecture:

The present invention can be employed in most organic EL device configurations. These include very simple structures comprising a single anode and cathode to more complex devices, such as passive matrix displays comprised
30 of orthogonal arrays of anodes and cathodes to form pixels, and active-matrix

displays where each pixel is controlled independently, for example, with thin film transistors (TFTs).

There are numerous configurations of the organic layers wherein the present invention can be successfully practiced. A typical structure is shown in FIG. 1 and includes a substrate 101, an anode 103, a hole-injecting layer 105, a hole-transporting layer 107, a light-emitting layer 109, an electron-transporting layer 111, and a cathode 113. These layers are described in detail below. This figure is for illustration only and the individual layer thickness is not scaled according to the actual thickness. Note that the substrate may alternatively be located adjacent to the cathode, or the substrate may actually constitute the anode or cathode. The organic layers between the anode and cathode are conveniently referred to as the organic EL element.

The anode 103 and cathode 113 of the OLED are connected to a voltage/current source 250 through electrical conductors 260. The OLED is operated by applying a potential between the anode 103 and cathode 113 such that the anode 103 is at a more positive potential than the cathode 113. Holes are injected into the organic EL element from the anode 103 and electrons are injected into the organic EL element at the anode 103. Enhanced device stability can sometimes be achieved when the OLED is operated in an AC mode where, for some time period in the cycle, the potential bias is reversed and no current flows. An example of an AC driven OLED is described in U.S. Patent No. 5,552,678. Substrate:

The OLED device of this invention is typically provided over a supporting substrate 101 where either the cathode 113 or anode 103 can be in contact with the substrate 101. The electrode in contact with the substrate 101 is conveniently referred to as the bottom electrode. Conventionally, the bottom electrode is the anode, but this invention is not limited to that configuration. The substrate 101 can either be light transmissive or opaque, depending on the intended direction of light emission. The light transmissive property is desirable for viewing the EL emission through the substrate 101. Transparent glass or plastic is commonly employed in such cases. The substrate 101 may be a

complex structure including multiple layers of materials. This is typically the case for active matrix substrates wherein TFTs are provided below the EL layers. It is still necessary that the substrate 101, at least in the emissive pixilated areas, be comprised of largely transparent materials such as glass or polymers. For applications where the EL emission is viewed through the top electrode, the transmissive characteristic of the bottom support is immaterial, and therefore can be light transmissive, light absorbing or light reflective. Substrates for use in this case include, but are not limited to, glass, plastic, semiconductor materials, silicon, ceramics, and circuit board materials. Again, the substrate 101 may be a complex structure including multiple layers of materials such as found in active matrix TFT designs. Of course it is necessary to provide in these device configurations a light-transparent top electrode.

Anode:

When EL emission is viewed through anode 103, the anode 103 should be transparent or substantially transparent to the emission of interest. Common transparent anode materials used in this invention are indium-tin oxide (ITO), indium-zinc oxide (IZO) and tin oxide, but other metal oxides can work including, but not limited to, aluminum- or indium-doped zinc oxide, magnesium-indium oxide, and nickel-tungsten oxide. In addition to these oxides, metal nitrides, such as gallium nitride, and metal selenides, such as zinc selenide, and metal sulfides, such as zinc sulfide, can be used as the anode 103. Anode 103 can be modified with plasma-deposited fluorocarbons as disclosed in EP Patent 0914025. For applications where EL emission is viewed only through the cathode electrode, the transmissive characteristics of anode are immaterial and any conductive material can be used, transparent, opaque or reflective. Example conductors for this application include, but are not limited to, gold, iridium, molybdenum, palladium, and platinum. Typical anode materials, transmissive or otherwise, have a work function of 4.1 eV or greater. Desired anode materials are commonly deposited by any suitable means such as evaporation, sputtering, chemical vapor deposition, or electrochemical means. Anodes can be patterned using well-known photolithographic processes. Optionally, anodes may be

polished prior to application of other layers to reduce surface roughness so as to minimize shorts or enhance reflectivity.

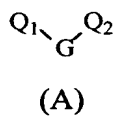
Hole-Injection Layer (HIL):

While not always necessary, it is often useful that a hole-injecting layer 105 be provided between anode 103 and hole-transporting layer 107. The hole-injecting material can serve to improve the film formation property of subsequent organic layers and to facilitate injection of holes into the hole-transporting layer 107. Suitable materials for use in the hole-injecting layer 105 include, but are not limited to, porphyrinic compounds as described in U.S. Patent No. 4,720,432, plasma-deposited fluorocarbon polymers as described in U.S. Patent No. 6,208,075, and some aromatic amines, for example, m-MTDATA (4,4',4''-tris[(3-methylphenyl)phenylamino]triphenylamine). Alternative hole-injecting materials reportedly useful in organic EL devices are described in EP 0 891 121 A1 and EP 1 029 909 A1.

Hole-Transporting Layer (HTL)

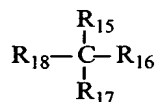
The hole-transporting layer 107 of the organic EL device in general contains at least one hole-transporting compound such as an aromatic tertiary amine, where the latter is understood to be a compound containing at least one trivalent nitrogen atom that is bonded only to carbon atoms, at least one of which is a member of an aromatic ring. In one form the aromatic tertiary amine can be an arylamine, such as a monoarylamine, diarylamine, triarylamine, or a polymeric arylamine. Exemplary monomeric triarylaminines are illustrated by Klupfel et al. U.S. Patent No. 3,180,730. Other suitable triarylaminines substituted with one or more vinyl radicals and/or including at least one active hydrogen containing group are disclosed by Brantley et al U.S. Patent Nos. 3,567,450 and 3,658,520.

A more preferred class of aromatic tertiary amines are those which include at least two aromatic tertiary amine moieties as described in U.S. Patent Nos. 4,720,432 and 5,061,569. Such compounds include those represented by structural formula (A).



wherein Q₁ and Q₂ are independently selected aromatic tertiary amine moieties and G is a linking group such as an arylene, cycloalkylene, or alkylene group of a carbon to carbon bond. In one embodiment, at least one of Q₁ or Q₂ contains a polycyclic fused ring structure, e.g., a naphthalene. When G is an aryl group, it is conveniently a phenylene, biphenylene, or naphthalene moiety.

A useful class of triarylaminines satisfying structural formula (A) and containing two triarylamine moieties is represented by structural formula (B):



10

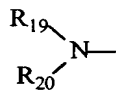
(B)

wherein:

R₁₅ and R₁₆ each independently represents a hydrogen atom, an aryl group, or an alkyl group or R₁₅ and R₁₆ together represent the atoms completing a cycloalkyl group; and

15

R₁₇ and R₁₈ each independently represents an aryl group, which is in turn substituted with a diaryl substituted amino group, as indicated by structural formula (C):



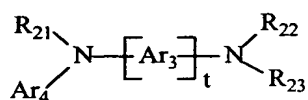
20

(C)

wherein R₁₉ and R₂₀ are independently selected aryl groups. In one embodiment, at least one of R₁₉ or R₂₀ contains a polycyclic fused ring structure, e.g., a naphthalene.

25

Another class of aromatic tertiary amines are the tetraaryldiamines. Desirable tetraaryldiamines include two diarylamino groups, such as indicated by formula (C), linked through an arylene group. Useful tetraaryldiamines include those represented by formula (D):



(D)

wherein

each Ar₃ is an independently selected arylene group, such as a phenylene or anthracene moiety,

t is an integer of from 1 to 4; and

Ar₄, R₂₁, R₂₂, and R₂₃ are independently selected aryl groups.

In a typical embodiment, at least one of Ar₄, R₂₁, R₂₂, and R₂₃ is a polycyclic fused ring structure, e.g., a naphthalene.

The various alkyl, alkylene, aryl, and arylene moieties of the foregoing structural formulae (A), (B), (C), (D), can each in turn be substituted. Typical substituents include alkyl groups, alkoxy groups, aryl groups, aryloxy groups, and halogen such as fluoride, chloride, and bromide. The various alkyl and alkylene moieties typically contain from about 1 to 6 carbon atoms. The cycloalkyl moieties can contain from 3 to about 10 carbon atoms, but typically contain five, six, or seven ring carbon atoms--eg, cyclopentyl, cyclohexyl, and cycloheptyl ring structures. The aryl and arylene moieties are usually phenyl and phenylene moieties.

The hole-transporting layer can be formed of a single or a mixture of aromatic tertiary amine compounds. Specifically, one may employ a triarylamine, such as a triarylamine satisfying the formula (B), in combination with a tetraaryldiamine, such as indicated by formula (D). When a triarylamine is employed in combination with a tetraaryldiamine, the latter is positioned as a layer interposed between the triarylamine and the electron injecting and transporting layer. Illustrative of useful aromatic tertiary amines are the following:

1,1-Bis(4-di-*p*-tolylaminophenyl)cyclohexane

1,1-Bis(4-di-*p*-tolylaminophenyl)-4-phenylcyclohexane

4,4'-Bis(diphenylamino)quadruphenyl

Bis(4-dimethylamino-2-methylphenyl)-phenylmethane

	N,N,N-Tri(<i>p</i> -tolyl)amine
	4-(di- <i>p</i> -tolylamino)-4'-[4(di- <i>p</i> -tolylamino)-styryl]stilbene
	N,N,N',N'-Tetra- <i>p</i> -tolyl-4,4'-diaminobiphenyl
	N,N,N',N'-Tetraphenyl-4,4'-diaminobiphenyl
5	N,N,N',N'-tetra-1-naphthyl-4,4'-diaminobiphenyl
	N,N,N',N'-tetra-2-naphthyl-4,4'-diaminobiphenyl
	N-Phenylcarbazole
	4,4'-Bis[N-(1-naphthyl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(1-naphthyl)-N-(2-naphthyl)amino]biphenyl
10	4,4''-Bis[N-(1-naphthyl)-N-phenylamino] <i>p</i> -terphenyl
	4,4'-Bis[N-(2-naphthyl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(3-acenaphthenyl)-N-phenylamino]biphenyl
	1,5-Bis[N-(1-naphthyl)-N-phenylamino]naphthalene
	4,4'-Bis[N-(9-anthryl)-N-phenylamino]biphenyl
15	4,4''-Bis[N-(1-anthryl)-N-phenylamino]- <i>p</i> -terphenyl
	4,4'-Bis[N-(2-phenanthryl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(8-fluoranthryl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(2-pyrenyl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(2-naphthacenyl)-N-phenylamino]biphenyl
20	4,4'-Bis[N-(2-perylenyl)-N-phenylamino]biphenyl
	4,4'-Bis[N-(1-coronenyl)-N-phenylamino]biphenyl
	2,6-Bis(di- <i>p</i> -tolylamino)naphthalene
	2,6-Bis[di-(1-naphthyl)amino]naphthalene
	2,6-Bis[N-(1-naphthyl)-N-(2-naphthyl)amino]naphthalene
25	N,N,N',N'-Tetra(2-naphthyl)-4,4''-diamino- <i>p</i> -terphenyl
	4,4'-Bis{N-phenyl-N-[4-(1-naphthyl)-phenyl]amino}biphenyl
	4,4'-Bis[N-phenyl-N-(2-pyrenyl)amino]biphenyl
	2,6-Bis[N,N-di(2-naphthyl)amine]fluorene
	1,5-Bis[N-(1-naphthyl)-N-phenylamino]naphthalene
30	4,4',4''-tris[(3-methylphenyl)phenylamino]triphenylamine

Another class of useful hole-transporting materials includes polycyclic aromatic compounds as described in EP 1 009 041. Tertiary aromatic amines with more than two amine groups may be used including oligomeric materials. In addition, polymeric hole-transporting/hole injection materials can be used such as poly(N-vinylcarbazole) (PVK), polythiophenes, polypyrrole, polyaniline (Yang, Y. et al. *Appl. Phys. Lett.* 1994, 64, 1245) and copolymers such as poly(3,4-ethylenedioxythiophene) / poly(4-styrenesulfonate) also called PEDOT/PSS (Groenendaal, L. B. et al. *Adv. Mater.* 2000, 12, 481).

Light-Emitting Layer (LEL)

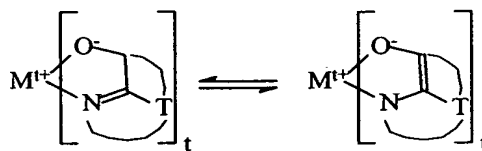
As more fully described in U.S. Patent Nos. 4,769,292 and 5,935,721, the light-emitting layer (LEL) 109 of the organic EL element includes a luminescent or fluorescent material where electroluminescence is produced as a result of electron-hole pair recombination in this region. The light-emitting layer 119 can include a single material including both small molecules and polymers. For small molecules, more commonly the LEL consists of a host material doped with a guest compound or compounds where light emission comes primarily from the dopant and can be of any color. The host materials in the light-emitting layer can be an electron-transporting material, as defined below, a hole-transporting material, as defined above, or another material or combination of materials that support hole-electron recombination. The dopant is usually chosen from highly fluorescent dyes, but phosphorescent compounds, e.g., transition metal complexes as described in WO 98/55561, WO 00/18851, WO 00/57676, and WO 00/70655 are also useful. Simultaneously, the color of the EL devices can be tuned using dopants of different emission wavelengths. By using a mixture of dopants, EL color characteristics of the combined spectra of the individual dopant are produced. This dopant scheme has been described in considerable detail for EL devices in U.S. Patent No. 4,769,292 for fluorescent dyes. Dopants are typically coated as 0.01 to 10 % by weight into the host material. Polymeric materials such as polyfluorenes and poly(arylene vinylenes) (e.g., poly(p-phenylenevinylene), PPV) can also be used as the host material. In this case, small molecule dopants

can be molecularly dispersed into the polymeric host, or the dopant could be added by copolymerizing a minor constituent into the host polymer.

An important relationship for choosing a dye as a dopant is a comparison of the bandgap potential which is defined as the energy difference between the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) of the molecule. For efficient energy transfer from the host to the dopant molecule, a necessary condition is that the band gap of the dopant is smaller than that of the host material. For phosphorescent emitters it is also important that the host triplet energy level of the host be high enough to enable energy transfer from host to dopant.

For small molecules, host and emitting molecules known to be of use include, but are not limited to, those disclosed in U.S. Patent Nos. 4,768,292; 5,141,671; 5,150,006; 5,151,629; 5,405,709; 5,484,922; 5,593,788; 5,645,948; 5,683,823; 5,755,999; 5,928,802; 5,935,720; 5,935,721, and 6,020,078.

For example, small molecule metal complexes of 8-hydroxyquinoline and similar derivatives (Formula E) constitute one class of useful host compounds capable of supporting electroluminescence, and are particularly suitable for light emission of wavelengths longer than 500 nm, e.g., green, yellow, orange, and red.



wherein:

M represents a metal;

t is an integer of from 1 to 4; and

T independently in each occurrence represents the atoms completing a nucleus having at least two fused aromatic rings.

From the foregoing it is apparent that the metal can be monovalent, divalent, trivalent, or tetravalent metal. The metal can, for example, be an alkali metal, such as lithium, sodium, or potassium; an alkaline earth metal, such as

magnesium or calcium; an earth metal, such aluminum or gallium, or a transition metal such as zinc or zirconium. Generally any monovalent, divalent, trivalent, or tetravalent metal known to be a useful chelating metal can be employed.

T completes a heterocyclic nucleus containing at least two fused aromatic rings, at least one of which is an azole or azine ring. Additional rings, including both aliphatic and aromatic rings, can be fused with the two required rings, if required. To avoid adding molecular bulk without improving on function the number of ring atoms is usually maintained at 18 or less.

Illustrative of useful chelated oxinoid compounds are the following:

CO-1: Aluminum trisoxine [alias, tris(8-quinolinolato)aluminum(III)]

CO-2: Magnesium bisoxine [alias, bis(8-quinolinolato)magnesium(II)]

CO-3: Bis[benzo{f}-8-quinolinolato]zinc (II)

CO-4: Bis(2-methyl-8-quinolinolato)aluminum(III)- μ -oxo-bis(2-methyl-8-quinolinolato) aluminum(III)

CO-5: Indium trisoxine [alias, tris(8-quinolinolato)indium]

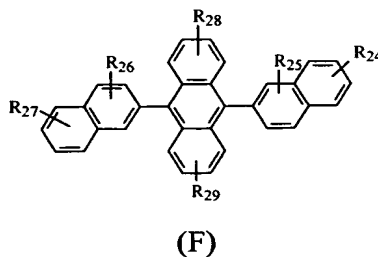
CO-6: Aluminum tris(5-methyloxine) [alias, tris(5-methyl-8-quinolinolato)aluminum(III)]

CO-7: Lithium oxine [alias, (8-quinolinolato)lithium(I)]

CO-8: Gallium oxine [alias, tris(8-quinolinolato)gallium(III)]

CO-9: Zirconium oxine [alias, tetra(8-quinolinolato)zirconium(IV)]

Derivatives of 9,10-di-(2-naphthyl)anthracene (Formula F) constitute one class of useful hosts capable of supporting electroluminescence, and are particularly suitable for light emission of wavelengths longer than 400 nm, e.g., blue, green, yellow, orange or red.



wherein: R₂₄, R₂₅, R₂₆, R₂₇, R₂₈, and R₂₉ represent one or more substituents on each ring where each substituent is individually selected from the following groups:

Group 1: hydrogen, or alkyl of from 1 to 24 carbon atoms;

5 Group 2: aryl or substituted aryl of from 5 to 20 carbon atoms;

Group 3: carbon atoms from 4 to 24 necessary to complete a fused aromatic ring of anthracenyl; pyrenyl, or perylenyl;

Group 4: heteroaryl or substituted heteroaryl of from 5 to 24 carbon atoms as necessary to complete a fused heteroaromatic ring of furyl, thienyl, pyridyl,
10 quinolinyl or other heterocyclic systems;

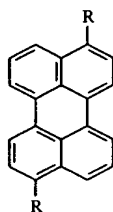
Group 5: alkoxyamino, alkylamino, or arylamino of from 1 to 24 carbon atoms; and

Group 6: fluorine, chlorine, bromine or cyano.

Illustrative examples include 9,10-di-(2-naphthyl)anthracene and 2-
15 *t*-butyl-9,10-di-(2-naphthyl)anthracene. Other anthracene derivatives can be useful as a host in the LEL, including derivatives of 9,10-bis[4-(2,2-diphenylethenyl)phenyl]anthracene.

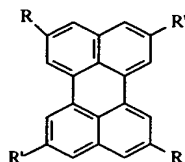
Distyrylarylene derivatives are also useful hosts, as described in U.S. Patent No. 5,121,029. Carbazole derivatives are particularly useful hosts for
20 phosphorescent emitters.

Useful fluorescent dopants (FD) include, but are not limited to, derivatives of anthracene, tetracene, xanthene, perylene, rubrene, coumarin, rhodamine, and quinacridone, dicyanomethylenepyran compounds, thiopyran compounds, polymethine compounds, pyrilium and thiapyrilium compounds, fluorene derivatives, perflanthene derivatives, indenoperylene derivatives, bis(azinyl)amine boron compounds, bis(azinyl)methane compounds, and carbostyryl compounds. Useful phosphorescent dopants (PD) include but are not limited to organometallic complexes of transition metals of iridium, platinum, palladium, or osmium. Illustrative examples of useful dopants include, but are not limited to, the following:



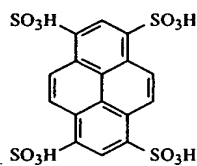
FD 1 R = H

FD 2 R = CO₂Pr-i

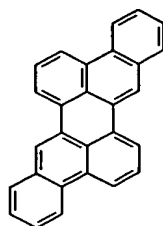


FD 3 R = H, R' = t-Bu

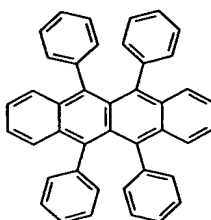
FD 4 R = R' = t-Bu



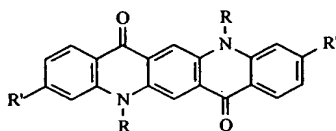
FD 5



FD 6



FD 7



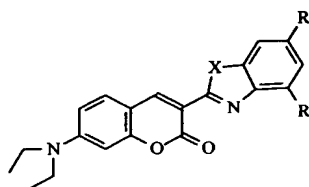
FD 8 $R = R' = H$

FD 9 $R = Me, R' = H$

FD 10 $R = Pr-i, R' = H$

FD 11 $R = Me, R' = F$

FD 12 $R = phenyl, R' = H$



FD 13 $R = R' = H, X = O$

FD 14 $R = H, R' = Me, X = O$

FD 15 $R = Me, R' = H, X = O$

FD 16 $R = Me, R' = Me, X = O$

FD 17 $R = H, R' = t-Bu, X = O$

5

FD 18 R = t-Bu, R' = H, X = O

FD 19 R = R' = t-Bu, X = O

FD 20 R = R' = H, X = S

FD 21 R = H, R' = Me, X = S

FD 22 R = Me, R' = H, X = S

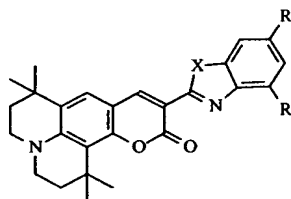
FD 23 R = Me, R' = Me, X = S

FD 24 R = H, R' = t-Bu, X = S

FD 25 R = t-Bu, R' = H, X = S

FD 26 R = R' = t-Bu, X = S

10



FD 27 R = R' = H, X = O

FD 28 R = H, R' = Me, X = O

FD 29 R = Me, R' = H, X = O

15

FD 30 R = Me, R' = Me, X = O

FD 31 R = H, R' = t-Bu, X = O

FD 32 R = t-Bu, R' = H, X = O

FD 33 R = R' = t-Bu, X = O

FD 34 R = R' = H, X = S

20

FD 35 R = H, R' = Me, X = S

FD 36 R = Me, R' = H, X = S

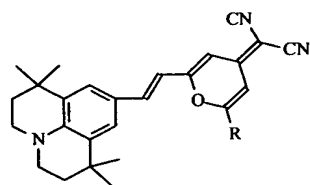
FD 37 R = Me, R' = Me, X = S

FD 38 R = H, R' = t-Bu, X = S

FD 39 R = t-Bu, R' = H, X = S

25

FD 40 R = R' = t-Bu, X = S



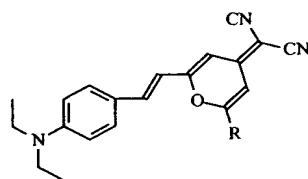
FD 41 R = phenyl

FD 42 R = Me

FD 43 R = t-Bu

FD 44 R = mesityl

5



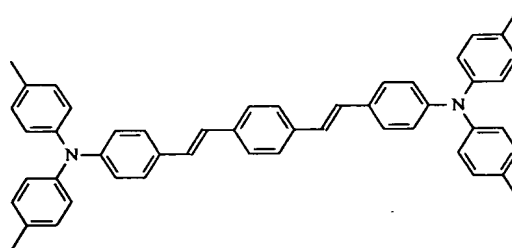
FD 45 R = phenyl

FD 46 R = Me

FD 47 R = t-Bu

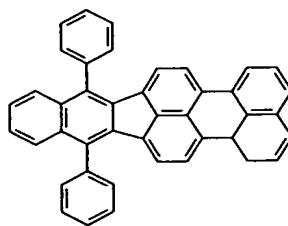
FD 48 R = mesityl

10

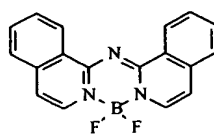


FD 49

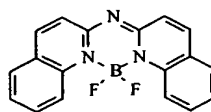
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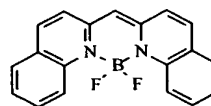
FD 50



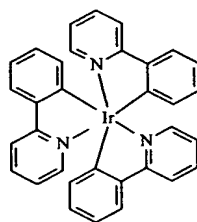
FD 51



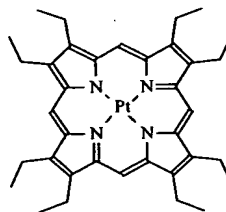
FD 52



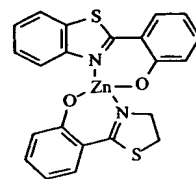
FD 53



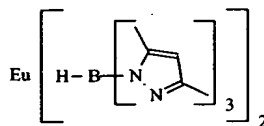
PD 1 (Ir(PPY)₃)



PD 2



PD 3



PD 4

5 Electron-Transporting Layer (ETL):

Preferred thin film-forming materials for use in forming the electron-transporting layer 111 of the organic EL devices of this invention are metal chelated oxinoid compounds, including chelates of oxine itself (also commonly referred to as 8-quinolinol or 8-hydroxyquinoline). Such compounds help to inject and transport electrons and exhibit both high levels of performance and are readily fabricated in the form of thin films. Exemplary of contemplated oxinoid compounds are those satisfying structural formula (E), previously described.

Other electron-transporting materials include various butadiene derivatives as disclosed in U.S. Patent No. 4,356,429 and various heterocyclic optical brighteners as described in U.S. Patent No. 4,539,507. Triazines are also known to be useful as electron transporting materials. Oxadiazole compounds including small molecules and polymers are useful electron transporting materials as described in U.S. Patent No. 6,451,457.

20 Cathode

When light emission is viewed solely through the anode, the cathode 113 used in this invention can include nearly any conductive material. Desirable materials have good film-forming properties to ensure good contact with the underlying organic layer, promote electron injection at low voltage, and have good stability. Useful cathode materials often contain a low work function metal (< 4.0 eV) or metal alloy. One preferred cathode material is comprised of a Mg:Ag alloy wherein the percentage of silver is in the range of 1 to 20 %, as described in U.S. Patent No. 4,885,221. Another suitable class of cathode materials includes bilayers comprising a thin electron-injection layer (EIL) in

contact with the organic layer (e.g., ETL) which is capped with a thicker layer of a conductive metal. Here, the EIL preferably includes a low work function metal or metal salt, and if so, the thicker capping layer does not need to have a low work function. One such cathode is comprised of a thin layer of LiF followed by a thicker layer of Al as described in U.S. Patent No. 5,677,572. Other useful cathode material sets include, but are not limited to, those disclosed in U.S. Patent Nos. 5,059,861; 5,059,862, and 6,140,763.

When light emission is viewed through the cathode, the cathode must be transparent or nearly transparent. For such applications, metals must be thin or one must use transparent conductive oxides, or a combination of these materials. Optically transparent cathodes have been described in more detail in U.S. Patent Nos. 4,885,211; 5,247,190; 5,703,436; 5,608,287; 5,837,391; 5,677,572; 5,776,622; 5,776,623; 5,714,838; 5,969,474; 5,739,545; 5,981,306; 6,137,223; 6,140,763; 6,172,459, 6,278,236, and 6,284,393; JP 3,234,963 and EP 1 076 368. Cathode materials are typically deposited by evaporation, sputtering, or chemical vapor deposition. When needed, patterning can be achieved through many well known methods including, but not limited to, through-mask deposition, integral shadow masking as described in U.S. Patent No. 5,276,380 and EP 0 732 868, laser ablation, and selective chemical vapor deposition.

Other Useful Organic Layers and Device Architecture

In some instances, layers 109 and 111 can optionally be collapsed into a single layer that serves the function of supporting both light emission and electron transportation, or layers 107 and 109 can optionally be collapsed into a single layer that serves the function of supporting both light emission and hole transportation. Alternatively, layers 107, 109 and 111 can optionally be collapsed into a single layer that serves the function of supporting both light emission and hole and electron transportation. This is the preferred EL device structure of this invention and is referred to as "single-layer" device.

It also known in the art that emitting dopants may be added to the hole-transporting layer, which may serve as a host. Multiple dopants may be added to one or more layers in order to create a white-emitting EL device, for

example, by combining blue- and yellow-emitting materials, cyan- and red-emitting materials, or red-, green-, and blue-emitting materials. White-emitting devices are described, for example, in EP 1 187 235, EP 1 182 244, U.S. Patent Application Publication No. 20020025419; U.S. Patent Nos. 5,683,823;
5 5,503,910; 5,405,709, and 5,283,182.

Additional layers such as electron or hole-blocking layers as taught in the art may be employed in devices of this invention. Hole-blocking layers are commonly used to improve efficiency of phosphorescent emitter devices, for example, as in U.S. Patent Application Publication No. 20020015859.

10 This invention may be used in so-called stacked device architecture, for example, as taught in U.S. Patent Nos. 5,703,436 and 6,337,492.
Deposition of organic layers

The organic materials mentioned above can be deposited as high quality transparent thin films by various methods such as a vapor deposition or
15 sublimation method, an electron-beam method, a sputtering method, a thermal transferring method, a molecular lamination method and a coating method such as solution casting, spin-coating or inkjet printing, with an optional binder to improve film formation. If the material is a polymer, solvent deposition is usually preferred. The material to be deposited by sublimation can be vaporized from a
20 sublimator "boat" often comprised of a tantalum material, e.g., as described in U.S. Patent No. 6,237,529, or can be first coated onto a donor sheet and then sublimed in closer proximity to the substrate. Layers with a mixture of materials can utilize separate sublimator boats or the materials can be pre-mixed and coated from a single boat or donor sheet. Patterned deposition can be achieved using
25 shadow masks, integral shadow masks (U.S. Patent No. 5,294,870), spatially-defined thermal dye transfer from a donor sheet (U.S. Patent Nos. 5,688,551; 5,851,709 and 6,066,357) and inkjet method (U.S. Patent No. 6,066,357).

Preferably, the spin-coating or inkjet printing technique is used to deposit the conjugated polymer of the invention, and only one polymer is
30 deposited in a single layer device.

Encapsulation:

Most organic EL devices are sensitive to moisture or oxygen, or both, so they are commonly sealed in an inert atmosphere such as nitrogen or argon, along with a desiccant such as alumina, bauxite, calcium sulfate, clays, silica gel, zeolites, alkaline metal oxides, alkaline earth metal oxides, sulfates, or metal halides and perchlorates. Methods for encapsulation and desiccation include, but are not limited to, those described in U.S. Patent No. 6,226,890. In addition, barrier layers such as SiO_x, Teflon, and alternating inorganic/polymeric layers are known in the art for encapsulation.

10 Optical Optimization:

Organic EL devices of this invention can employ various well-known optical effects in order to enhance its properties if desired. This includes optimizing layer thicknesses to yield maximum light transmission, providing dielectric mirror structures, replacing reflective electrodes with light-absorbing electrodes, providing anti glare or anti-reflection coatings over the display, providing a polarizing medium over the display, or providing colored, neutral density, or color conversion filters over the display. Filters, polarizers, and anti-glare or anti-reflection coatings may be specifically provided over the cover or as part of the cover.

20

EXAMPLES

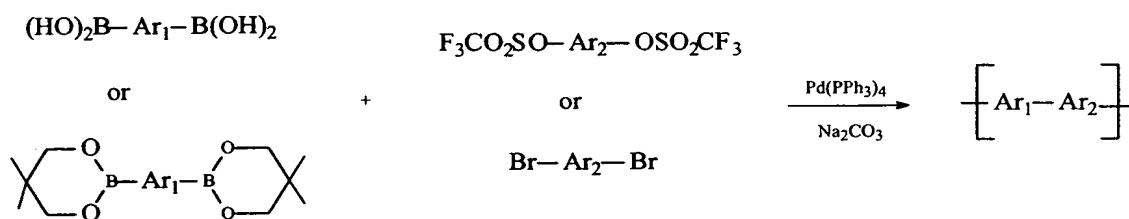
The invention and its advantages are further illustrated by the following specific examples:

Synthesis of Monomers

The monomers to be used in the present invention to construct polymers are not necessary to be particularly restricted. Any monomers can be used as long as the polymer formed is a polymer which satisfies the general formulas (II) and (III). Typical synthesis is illustrated in Schemes 1-4.

25





Scheme 4

5 Example 1: synthesis of compound 1

To a 500 mL round-bottom flask were added 2-fluoronitrobenzene (16.0 g, 0.11 mol), 4-hexylaniline (23.7 g, 0.13 mol), tripropylamine (20.7 g, 0.14 mol), and 150 mL of NMP. The reaction was heated to 120 °C overnight under nitrogen. After cooling down to room temperature, the reaction mixture was extracted with methylene chloride and water, and the combined organic phase was washed 3 times with water. The crude product was obtained as brown oil and was purified by column chromatography on silica gel using petroleum ether as an eluent to give 31.4 g of pure product as orange-red oil (93% yield). ¹H NMR (CDCl₃) δ (ppm): 0.90 (t, J = 6.8 Hz, 3 H), 1.26-1.38 (m, 6 H), 1.60-1.65 (m, 2 H), 2.62 (t, J = 7.7 Hz, 2 H), 6.70-6.76 (m, 1 H), 7.15-7.25 (m, 5 H), 7.31-7.58 (m, 1 H), 8.19 (dd, J₁ = 8.6 Hz, J₂ = 1.4 Hz, 1 H), 9.46 (s, br, 1 H); ¹³C NMR (CDCl₃) δ (ppm): 14.08, 22.59, 28.94, 31.42, 31.69, 35.45, 115.96, 117.04, 124.66, 126.58, 129.60, 132.80, 135.61, 136.031, 140.78, 143.62; FD-MS: *m/z* 298 (M⁺).

Example 2: synthesis of compound 2

Compound 1 (17.5 g, 0.059 mol) and sodium sulfide nonahydrate (42.0 g, 0.18 mol) were added to 150 mL of ethanol. The reaction mixture was heated to reflux for 9 hours. After the removal of the solvent, the residue was extracted with water and methylene chloride. The crude product was recrystallized from petroleum ether to give 13.3 g of white flaky crystals (84% yield). ¹H NMR (CDCl₃) δ (ppm): 0.88 (t, J = 6.1 Hz, 3 H), 1.29-1.56 (m, 8 H), 2.51 (t, J = 7.6 Hz, 2 H), 3.64 (s, br, 2 H), 5.07 (s, br, 1 H), 6.65-6.77 (m, 4 H), 6.94-7.09 (m, 4 H); ¹³C NMR (CDCl₃) δ (ppm): 14.21, 22.70, 29.07, 31.80,

35.17, .53, 116.00, 119.03, 123.78, 124.94, 129.00, 129.25, 133.95, 141.19, 142.65; M.p. 66-67 °C; FD-MS: m/z 268 (M^+).

Example 3: synthesis of compound 3

2,5-Dibromobenzoic acid (15.0 g, 0.054 mol) was heated to reflux
5 in 20 mL of thionyl chloride (32.7 g, 0.28 mol) overnight. The solid dissolved slowly upon heating and the reaction turned into a light yellow solution. Excess of thionyl chloride was distilled off and the solid residue was recrystallized from dry hexane to give 14.1 g of pure product as off-white solid (88% yield). ^1H NMR (CDCl_3) δ (ppm): 7.55 (d, $J = 2.1$ Hz, 1 H), 7.56 (s, 1 H), 8.13 (d, $J = 2.1$
10 Hz, 1 H); ^{13}C NMR (CDCl_3) δ (ppm): 119.86, 121.12, 135.42, 135.92, 136.26, 137.04; M.p. 119-120 °C.

Example 4: synthesis of compound 4

Compound 2 (5.0 g, 0.019 mol) was dissolved in 50 mL of methylene chloride and cooled to 0 °C. To the solution was added triethylamine
15 (2.0 g, 0.020 mol) and compound 3 (5.6 g, 0.019 mol). The reaction was stirred at room temperature overnight, extracted with water, and dried over MgSO_4 . The crude product was obtained as 9.5 g of off-white solid (96% yield) and was used without purification. ^1H NMR (CDCl_3) δ (ppm): 0.88 (t, $J = 6.7$ Hz, 3 H), 1.26-1.36 (m, 6 H), 1.51-1.59 (m, 2 H), 2.52 (t, $J = 7.7$ Hz, 2 H), 5.54 (s, br, 1 H), 6.72
20 (d, $J = 8.3$ Hz, 2 H), 7.03 (d, $J = 8.3$ Hz, 2 H), 7.13-7.16 (m, 2 H), 7.22-7.25 (m, 1 H), 7.33-7.42 (m, 2 H), 7.46 (d, $J = 2.1$ Hz, 1 H), 8.02-8.06 (m, 1 H), 8.13 (s, br, 1 H); ^{13}C NMR (CDCl_3) δ (ppm): 14.09, 22.60, 28.97, 31.62, 31.72, 35.15, 116.80, 118.08, 121.52, 122.84, 123.35, 124.43, 126.20, 129.27, 130.89, 132.32, 134.42, 134.65, 134.81, 135.50, 139.13, 142.13, 164.36; M.p. 120-121 °C; FD-MS: m/z
25 530 (M^+).

Example 5: synthesis of compound 5

Compound 4 (9.2 g, 0.017 mol) was heated to reflux in 10 mL of POCl_3 (16.5 g, 0.11 mol) overnight. Excess POCl_3 was distilled off and the
30 residue was extracted with ethyl acetate and water. The crude product was obtained as a light brown foam and was purified by column chromatography on

silica gel using ethyl acetate/petroleum ether (8/92 v/v) as an eluent to give 7.2 g of pure product as viscous oil that solidified upon standing (81% yield). ^1H NMR (CDCl_3) δ (ppm): 0.88 (t, $J = 6.7$ Hz, 3 H), 1.28-1.34 (m, 6 H), 1.57-1.64 (m, 2 H), 2.62 (t, $J = 7.8$ Hz, 2 H), 7.14-7.39 (m, 9 H), 7.63 (s, 1 H), 7.91 ($J = 7.8$ Hz, 1 H); ^{13}C NMR (CDCl_3) δ (ppm): 14.02, 22.50, 28.79, 30.99, 31.56, 35.45, 110.76, 120.17, 120.78, 122.41, 123.12, 123.85, 126.34, 129.34, 132.98, 134.07, 134.09, 135.25, 135.64, 142.33, 143.35, 149.98; M.p. 64-65 °C; FD-MS: m/z 512 (M^+).

Example 6: synthesis of compound 6

2,5-Dibromo-p-xylene (35 g, 0.13 mol) and potassium permanganate (210 g, 1.32 mol) were mixed in 1.2 L of water. The reaction was heated to reflux for 2 days until the purple color of permanganate disappeared. The reaction was cooled and filtered through a pad of Celite. The filtrate was cooled in an ice-bath and neutralized by concentrated HCl solution until pH ~ 1. The white crystalline product was filtered, washed with water, and dried over P_2O_5 . ^1H NMR (DMSO) δ (ppm): 5.06 (s, br, 2 H), 8.02 (s, 2 H); M.p. 178-180 °C; FD-MS: m/z 324 (M^+).

Example 7: synthesis of compound 7

Compound 6 (19.2 g, 0.059 mol) was suspended in 30 mL of thionyl chloride and heated to reflux. Large amount of gas evolved in first 20 min. and the reaction became clear. The reaction was refluxed for 4 h and the excess of thionyl chloride was distilled off. To the light yellow residue was added dry hexane and the mixture was heated to reflux until all solid dissolved. The solution was cooled down and white crystals formed. The crystals were filtered off, washed with dry hexane, and dried under vacuum at room temperature overnight to give 19.2 g product as white solid at 90% yield. ^1H NMR (CDCl_3) δ (ppm): 8.21 (s, 2 H); ^{13}C NMR (CDCl_3) δ (ppm): 119.30, 137.10, 139.64, 164.28; Mp 80-82°C.

Example 8: synthesis of compound 8

Compound 2 (7.5 g, 0.028 mol) was dissolved in 150 mL of methylene chloride and cooled to 0 °C. To the solution was added triethylamine

(3.1 g, 0.031 mol) and compound 7 (5.0 g, 0.014 mol). The reaction was stirred at room temperature overnight. Large amount of precipitate formed. The precipitate was filtered and washed with water and ethyl acetate. The filtrate was extracted with ethyl acetate, and dried over MgSO_4 . Solvent was removed and the residue was combined with the yellow precipitate to give total 8.5 g of crude product (74% yield) and was used without purification. ^1H NMR (DMSO) δ (ppm): 0.84 (t, $J = 6.7$ Hz, 6 H), 1.26-1.36 (m, 16 H), 1.52-1.60 (m, 4 H), 6.82 (d, $J = 8.3$ Hz, 4 H), 6.95-6.97 (m, 2 H), 7.03 (d, $J = 8.3$ Hz, 4 H), 7.13-7.16 (m, 2 H), 7.22-7.23 (m, 2 H), 7.56 (d, $J = 2.1$ Hz, 2 H), 7.66 (s, 2 H), 10.01 (s, 1 H); ^{13}C NMR (DMSO) δ (ppm): 13.94, 22.07, 28.33, 31.06, 31.14, 34.48, 117.03, 118.22, 119.34, 120.89, 126.32, 126.44, 127.63, 128.82, 132.55, 132.61, 133.78, 137.42, 141.39, 164.27; M.p. 239-240 $^\circ\text{C}$; FD-MS: m/z 824 (M^+).

Example 9: synthesis of compound 9

Compound 8 (8.3 g, 0.010 mol) was heated to reflux in 20 mL of POCl_3 (32.9 g, 0.22 mol) overnight. The starting material did not dissolve completely and after overnight refluxing, the reaction turned into brown purple with gray precipitate. Excess POCl_3 was distilled off and the residue was suspended in ethyl acetate and water. The precipitate was filtered and washed with ethyl acetate to give 8.5 g of crude product as white powder. The crude product was purified by column chromatography on silica gel using ethyl acetate/methylene chloride (10/90 v/v) as an eluent to give 7.0 g of pure product as off-white powder (88% yield). ^1H NMR (CDCl_3) δ (ppm): 0.90 (t, $J = 6.8$ Hz, 6 H), 1.26-1.40 (m, 12 H), 1.62-1.69 (m, 4 H), 2.66 (t, $J = 7.8$ Hz, 4 H), 7.14 (d, $J = 8.3$ Hz, 4 H), 7.23 (d, $J = 8.3$ Hz, 4 H), 7.32-7.40 (m, 6 H), 7.67 (s, 2 H), 7.88 (d, $J = 8.2$ Hz, 2 H); ^{13}C NMR (CDCl_3) δ (ppm): 11.97, 20.16, 26.55, 28.83, 29.20, 32.97, 108.74, 117.78, 120.90, 121.86, 124.46, 127.28, 130.66, 132.50, 133.12, 133.47, 134.18, 140.36, 140.88; M.p. 215-217 $^\circ\text{C}$; FD-MS: m/z 788 (M^+).

Example 10: synthesis of compound 10

2-Nitroaniline (12.2 g, 0.089 mol) and 2-fluoro-1,4-dibromobenzene (33.7 g, 0.13 mol) was dissolved in 100 mL of DMF. To this

solution was added sodium bicarbonate (14.0 g, 0.17 mol). The reaction was degassed with nitrogen and heated to 150 °C for 2 days. Large amount of 2-nitroaniline was still present, but no 2-fluoro-1,4-dibromobenzene was observed on TLC. After cooling down, the reaction was extracted with water and
5 methylene chloride, and the combined organic phase was back-washed with water 3 times. The crude product was purified by column chromatography on silica gel using petroleum ether as an eluent and then recrystallized from heptane to give 1.5 g of pure product as orange solid (4.5 % yield). ¹H NMR (CDCl₃) δ (ppm): 6.91-6.96 (m, 1 H), 7.17 (dd, J₁ = 8.5 Hz, J₂ = 2.2 Hz, 1 H), 7.25-7.28 (m, 1 H), 7.46-
10 7.54 (m, 2 H), 7.59 (d, J = 2.2 Hz, 1 H), 8.23 (dd, J₁ = 8.5 Hz, J₂ = 1.3 Hz, 1 H); ¹³C NMR (CDCl₃) δ (ppm): 116.56, 116.86, 119.45, 119.52, 121.48, 125.77, 126.81, 128.68, 128.78, 134.70, 135.71, 140.18; M.p. 166-168 °C; FD-MS: *m/z* 370 (M⁺).

Example 11: synthesis of compound 11

15 Compound 10 (1.3 g, 3.5 mmol) and sodium sulfide nonahydrate (2.5 g, 0.10 mol) were added to 10 mL of ethanol. The reaction mixture was heated to reflux for 1 hours and then poured into cold water. The precipitate was filtered and washed with water and dried to give 0.80 g of crude product as light yellow solid (68% yield). ¹H NMR (CDCl₃) δ (ppm): 3.80 (s, br, 2 H), 5.72 (s, br,
20 1 H), 6.64 (d, J = 2.1 Hz, 1 H), 6.75-6.84 (m, 3 H), 7.08-7.14 (m, 2 H), 7.32 (d, J = 8.4 Hz, 1 H); ¹³C NMR (CDCl₃) δ (ppm): 116.22, 116.44, 119.10, 122.18, 122.23, 127.45, 127.73, 133.46; M.p. 106-107 °C; FD-MS: *m/z* 340 (M⁺).

Example 12: synthesis of compound 12

25 Compound 11 (0.60 g, 1.8 mmol) was dissolved in 10 mL of methylene chloride and cooled to 0 °C. To the solution was added triethylamine (0.21 g, 2.1 mmol) and 4-hexylbenzoyl chloride (0.39 g, 1.8 mmol). The reaction was stirred at room temperature for 2 hours. Some precipitate formed. The reaction was extracted with water, and dried over MgSO₄. Solvent was removed and the residue was recrystallized from heptane to give 0.68 g off-white crystals.
30 ¹H NMR (CDCl₃) δ (ppm): 0.93 (t, J = 6.6 Hz, 3 H), 1.31-1.40 (m, 6 H), 1.61-1.68

(m, 2 H), 2.69 (t, $J = 7.7$ Hz, 2 H), 6.23 (s, br, 1 H), 6.82 (d, $J = 2.2$ Hz, 1 H), 6.87 (dd, $J_1 = 8.4$ Hz, $J_2 = 2.2$ Hz, 1 H), 7.23-7.31 (m, 3 H), 7.35 (d, $J = 7.5$ Hz, 2 H), 7.40 (d, $J = 8.3$ Hz, 1 H), 7.72 (d, $J = 8.2$ Hz, 2 H), 8.26 (d, $J = 8.0$ Hz, 1 H), 8.34 (s, br 1 H); ^{13}C NMR (CDCl_3) δ (ppm): 14.06, 22.55, 28.90, 31.10, 31.63, 35.84, 109.73, 117.26, 122.52, 123.33, 125.54, 126.35, 127.10, 128.83, 131.07, 133.78, 133.99, 147.57, 165.79; M.p. 128-129 °C; FD-MS: m/z 528 (M^+).

Example 13 :synthesis of compound 13

Compound 12 (0.68 g, 1.3 mmol) was refluxed in 0.6 mL of POCl_3 (1.0 g, 6.4 mmol) for 30 minutes. Excess POCl_3 was distilled off and the residue was extracted with ethyl acetate and water. The crude product was purified by column chromatography on silica gel using ethyl acetate/petroleum ether (10/90 v/v) as an eluent to give 0.5 g of pure product as light greenish solid (77% yield). ^1H NMR (CDCl_3) δ (ppm): 0.87 (t, $J = 6.6$ Hz, 3 H), 1.26-1.32 (m, 6 H), 1.55-1.62 (m, 2 H), 2.60 (t, $J = 7.6$ Hz, 2 H), 7.00 (d, $J = 7.9$ Hz, 1 H), 7.14 (d, $J = 8.1$ Hz, 2 H), 7.24-7.29 (m, 1 H), 7.32-7.37 (m, 1 H), 7.49-7.53 (m, 4 H), 7.63 (d, $J = 8.2$ Hz, 1 H), 7.88 (d, $J = 7.9$ Hz, 1 H); ^{13}C NMR (CDCl_3) δ (ppm): 14.05, 22.55, 28.84, 30.95, 31.62, 35.73, 110.29, 119.90, 121.73, 123.20, 123.32, 123.44, 128.45, 128.50, 128.62, 128.81, 133.31, 133.82, 135.11, 145.01; M.p.156-158 °C; FD-MS: m/z 510 (M^+)

Synthesis of Polymers

Example 14: general procedure for synthesis of polymers via the Suzuki coupling reaction

Equal molar of aromatic di-bromide or di-triflate and aromatic di-boron compound, and phase transfer reagent Aliquat® 336 (0.10 equivalent to monomer) were dissolved in of toluene (the ratio of toluene to water (v/v) is about 3/1). To this solution was added 2 M Na_2CO_3 aqueous solution (3.3 equivalent to monomer). The reaction mixture was bubbled with dry nitrogen for 15 min and catalyst tetrakis(triphenylphosphine)palladium (0.03 equivalent to monomer) was added. The reaction was heated under vigorous reflux for 12-24 h, and small amount of phenylboronic acid was added for end-capping of bromo or triflate

group. The reaction was heated for 5 h and bromobenzene was added to end-cap boron group. The reaction was heated for another 4 h and then poured into methanol. The precipitated polymer was washed with methanol, diluted HCl solution, and dried. The polymer was treated with diethyl dithiocarbamate twice:
5 polymer was dissolved in toluene, and sodium diethyl dithiocarbamate in water (1 g in 10 mL of water) was added, and the mixture was stirred under nitrogen at 60 °C overnight. The toluene layer was separated, washed with water twice, and concentrated, and the polymer was precipitated into methanol twice. Polymer can then be extracted with acetone with a Soxhlet setup overnight to remove
10 oligomers. Polymer was dried under vacuum at 45 °C.

EL Device Fabrication and Performance

Example 15

An EL device satisfying the requirements of the invention was constructed in the following manner. The organic EL medium has a single layer
15 of the organic compound described in this invention.

- a) An indium-tin-oxide (ITO) coated glass substrate was sequentially ultra-sonicated in a commercial detergent, rinsed with deionized water, degreased in toluene vapor and exposed to ultraviolet light and ozone for a few minutes.
- 20 b) An aqueous solution of PEDOT (1.3% in water, Baytron P Trial Product AI 4083 from H. C. Stark) was spin-coated onto ITO under a controlled spinning speed to obtain thickness of 500 Angstroms. The coating was baked in an oven at 110 °C for 10 min.
- 25 c) A toluene solution of a polymer (300 mg in 30 mL of solvent), or a polymer with dopant(s) (dopant(s) are 4 wt% to polymer, for example, PD 1 Ir(PPY)₃ was used) was filtered through a 0.2 µm Teflon filter. The solution was then spin-coated onto PEDOT under a controlled spinning speed. The thickness of the film was between 500-1000 Angstroms.

- d) On the top of the organic thin film was deposited a cathode layer consisting of 15 angstroms of a CsF salt, followed by a 2000 angstroms of a 10:1 atomic ratio of Mg and Ag.

The above sequence completed the deposition of the EL device.

- 5 The device was then hermetically packaged in a dry glove box for protection against ambient environment.

- Table 1 summarizes the characterization of the polymers prepared in the present invention. Absorption (AB) and photoluminescence (PL) spectra were obtained from solid thin films of the polymers and EL spectra were obtained from ITO/PEDOT/organic material/CsF/Mg:Ag EL devices. The fabrication of EL devices was illustrated in Example 15. FIG. 2 shows the AB and PL spectra of polymer 43 in dilute toluene solution and thin film. FIG. 3 shows the EL spectra of devices from polymer 43 and polymer 43 doped with Ir(PPY)₃: ITO/PEDOT/polymer 43/CsF/Mg:Ag and ITO/PEDOT/polymer 43 + 4 wt% Ir(PPY)₃/CsF/Mg:Ag EL. And FIG. 4 shows the voltage-current characteristics of the doped and un-doped EL devices of polymer 43.

Table 1. Characterization of polymers according to Examples.

Polymer	M _w ^a	PDI	T _d (°C)	T _g (°C)	AB ^b (λ _{max} nm)	PL ^c (λ _{max} nm)	EL (λ _{max} nm)
5	NA ^d	NA	453	141	340	412 (342)	500
12	NA	NA	453	153	308	410 (310)	NA
24	4740	1.33	433	160	326	406 (332)	NA
43	4930	1.26	466	171	320	474 (380)	472
75	8470	1.42	492	155	378	428 (380)	460
90	3510	1.31	455	136	304	438 (306)	452
102	2570	1.41	452	131	302	442 (304)	456

- ^a weight average molecular weight, determined by size exclusion chromatography in THF using polystyrene standard. ^b as solid state thin film ^c as solid state thin film, the number in the parenthesis is the excitation wavelength ^d not available.

The entire contents of the patents and other publications referred to in this specification are incorporated herein by reference.

The invention has been described in detail with particular reference to certain preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

PARTS LIST

- 101 Substrate
- 103 Anode
- 105 Hole-Injecting layer (HIL)
- 107 Hole-Transporting layer (HTL)
- 109 Light-Emitting layer (LEL)
- 111 Electron-Transporting layer (ETL)
- 113 Cathode
- 250 Current / Voltage source
- 260 Electrical conductors